



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<b>(21) International Application Number:</b> PCT/CA92/00211 <b>(22) International Filing Date:</b> 20 May 1992 (20.05.92) <b>(30) Priority data:</b> 703,379 21 May 1991 (21.05.91) US <b>(60) Parent Application or Grant</b> (63) Related by Continuation US 703,379 (CIP) Filed on 21 May 1991 (21.05.91) <b>(71) Applicant (for all designated States except US):</b> BIOCHEM PHARMA INC. [CA/CA]; 2550 Daniel Johnson Boulevard, Suite 600, Laval, Quebec H7T 2L1 (CA). <b>(72) Inventors; and</b> <b>(75) Inventors/Applicants (for US only) :</b> MANSOUR, Tarek [CA/CA]; JIN, Haolun [CN/US]; TSE, Allan, H., L. [GB/CA]; SIDDIQUI, M., Arshad [IN/CA]; 531 des Prairies Boulevard, Building 10, Laval, Quebec H7V 1B7 (CA).	<b>(74) Agents:</b> MORROW, Joy, D. et al.; Smart & Biggar, 900-55 Metcalfe Street, P.O. Box 2999, Station D, Ottawa, Ontario K1P 5Y6 (CA).  <b>(81) Designated States:</b> AT, AT (European patent), AU, BB, BE (European patent), BF (OAPI patent), BG, BJ (OAPI patent), BR, CA, CF (OAPI patent), CG (OAPI patent), CH, CH (European patent), CI (OAPI patent), CM (OAPI patent), CS, DE, DE (European patent), DK, DK (European patent), ES, ES (European patent), FI, FR (European patent), GA (OAPI patent), GB, GB (European patent), GN (OAPI patent), GR (European patent), HU, IT (European patent), JP, KP, KR, LK, LU, LU (European patent), MC (European patent), MG, ML (OAPI patent), MN, MR (OAPI patent), MW, NL, NL (European patent), NO, PL, RO, RU, SD, SE, SE (European patent), SN (OAPI patent), TD (OAPI patent), TG (OAPI patent), US.  <b>Published</b> <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>	
<b>(54) Title:</b> PROCESSES FOR THE DIASTEREOSELECTIVE SYNTHESIS OF NUCLEOSIDES  <b>(57) Abstract</b>  The present invention relates to highly diastereoselective processes for production of <i>cis</i> -nucleosides and nucleoside analogues and derivatives in high optical purity and intermediates useful in those processes.		

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PROCESSES FOR THE DIASTEREOSELECTIVE  
SYNTHESIS OF NUCLEOSIDES

FIELD OF THE INVENTION

5           The present invention relates to  
diastereoselective processes for preparing optically  
active *cis*-nucleosides and nucleoside analogues and  
derivatives. The novel processes of this invention  
allow the stereo-controlled synthesis of a given  
10 enantiomer of a desired *cis*-nucleoside or nucleoside  
analogue or derivative in high optical purity. This  
invention also relates to novel intermediates useful in  
the processes of this invention.

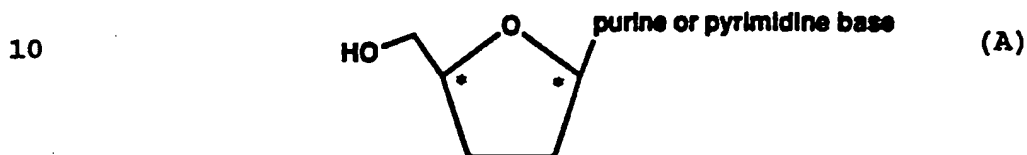
BACKGROUND OF THE INVENTION

15           Nucleosides and their analogues and  
derivatives are an important class of therapeutic  
agents. For example, a number of nucleosides have  
shown antiviral activity against retroviruses such as  
human immunodeficiency virus (HIV), hepatitis B virus  
20 (HBV) and human T-lymphotropic virus (HTLV) (PCT  
publication WO 89/04662 and European Patent publication  
0349242 A2). Among the nucleosides shown to have  
antiviral activity are 3'-azido-3'-deoxythymidine  
(AZT), 2'3'-dideoxy-cytidine (DDC), 2-hydroxymethyl-5-  
25 (cytosin-1'-yl)-1,3-oxathiolane and 2-hydroxymethyl-4-  
(guanine-9'-yl)-1,3-dioxolane (European Patent

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publication 0382526 A2 and European Patent publication 0377713 A2).

Most nucleosides and nucleoside analogues and derivatives contain at least two chiral centers (shown as \* in formula (A)), and exist in the form of two pairs of optical isomers (i.e., two in the *cis*-configuration and two in the *trans*- configuration). However, generally only the *cis*-isomers exhibit useful biological activity.



Different enantiomeric forms of the same *cis*-nucleoside may, however, have very different antiviral activities. M.M. Mansuri et al., "Preparation Of The Geometric Isomers Of DDC, DDA, D4C and D4T As Potential Anti-HIV Agents", Bioorg.Med.Chem. Lett., 1 (1), pp. 65-68 (1991). Therefore, a general and economically attractive stereoselective synthesis of the enantiomers of the biologically active *cis*-nucleosides is an important goal.

20 Many of the known processes for producing optically active nucleosides and their analogues and derivatives modify naturally occurring (i.e., optically active) nucleosides by altering the base or by altering the sugar via reductive procedures such as

25 deoxygenation or radical initiated reductions. C.K. Chu et al., "General Synthesis Of 2',3'-Dideoxynucleosides And 2',3'-Didehydro-2',3'-Dideoxynucleosides," J.Org.Chem., 54, pp. 2217-2225 (1989). These transformations involve multiple steps,

30 including protection and deprotection and usually

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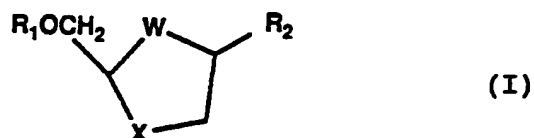
result in low yields. Moreover, they begin with and maintain the optical activity of the starting nucleoside. Thus, the nucleosides produced by these processes are limited to specific analogues of the enantiomeric form of the naturally occurring nucleoside. In addition, these procedures require the availability of the naturally occurring nucleoside, often an expensive starting material.

Other known processes for producing optically active nucleosides rely on conventional glycosylation procedures to add the sugar to the base. These procedures invariably give anomeric mixtures of *cis*- and *trans*-isomers which require tedious separation and result in lower yields of the desired biologically active *cis*-nucleoside. Improved glycosylation methods designed to yield only the *cis*-nucleoside require addition of a 2'- or 3'-substituent to the sugar. Because the 2'- or 3'-substituent is only useful in controlling *cis*-nucleoside synthesis in one configuration (when the 2' or 3' substituent is *trans*- to the 4' substituent), multiple steps are required to introduce this substituent in the proper configuration. The 2'- or 3'-substituent must then be removed after glycosylation, requiring additional steps. L. Wilson and D. Liotta, "A General Method For Controlling Stereochemistry In The Synthesis Of 2'-Deoxyribose Nucleosides", Tetrahedron Lett., 31, pp. 1815-1818 (1990). Furthermore, to obtain an optically pure nucleoside product, the starting sugar must be optically pure. This also requires a series of time-consuming syntheses and purification steps.

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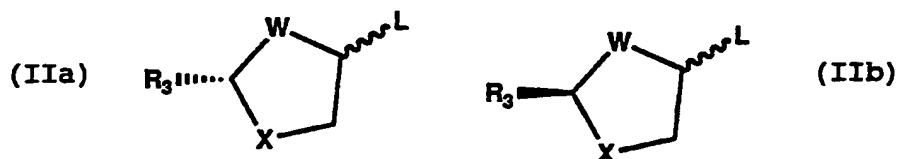
SUMMARY OF THE INVENTION

The present invention overcomes the difficulties and shortcomings of the prior art and provides processes for producing optically active *cis*-  
 5 nucleosides (1,3-oxathiolanes, 2,4-dioxolanes, and 1,3-dithiolanes) or nucleoside analogues and derivatives of formula (I)



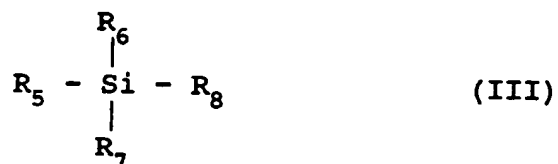
- wherein W is S, S=O, SO<sub>2</sub>, or O;  
 10 X is S, S=O, SO<sub>2</sub>, or O;  
 R<sub>1</sub> is hydrogen or acyl; and  
 R<sub>2</sub> is a purine or pyrimidine base or an analogue or derivative thereof.

The processes of this invention comprise the  
 15 step of glycosylating a desired purine or pyrimidine base or analogue or derivative thereof with an intermediate of formula (IIa) or (IIb)



wherein R<sub>3</sub> is a substituted carbonyl or carbonyl  
 20 derivative and L is a leaving group. Glycosylation is accomplished using a Lewis acid of the formula (III)

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wherein  $R_5$ ,  $R_6$ ,  $R_7$ , and  $R_8$  are defined below and the resulting intermediate is reduced to give a nucleoside or nucleoside analogue or derivative of formula (I).

The processes of this invention have the advantages of allowing preparation of a nucleoside of formula (I) (or analogues or derivatives thereof) without using expensive starting materials, cumbersome protection and deprotection steps or addition and removal of 2'- or 3'-substituents. The processes of this invention produce nucleosides in high yields, with high purity and high optical specificity. The processes of this invention have the further advantage of generating nucleosides whose stereoisomeric configuration can be easily controlled simply by the selection of the appropriate starting materials.

#### DETAILED DESCRIPTION OF THE INVENTION

In the processes for preparing optically active compounds of this invention in a configurational- and diastereo-selective manner, the following definitions are used:

$R_2$  is a purine or pyrimidine base or an analogue or derivative thereof.

A purine or pyrimidine base is a purine or pyrimidine base found in naturally occurring nucleosides. An analogue thereof is a base which mimics such naturally occurring bases in that their structures (the kinds of atoms and their arrangement) are similar to the naturally occurring bases but may either possess additional or lack certain of the functional properties of the naturally occurring bases.

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Such analogues include those derived by replacement of a CH moiety by a nitrogen atom, e.g., 5-azapyrimidines such as 5-azacytosine) or vice versa (e.g., 7-deazapurines, such as 7-deazaadenine or 7-deazaguanine) or both (e.g., 7-deaza, 8-azapurines). By derivatives of such bases or analogues are meant those bases wherein ring substituents are either incorporated, removed, or modified by conventional substituents known in the art, e.g., halogen, hydroxyl, amino, C<sub>1-6</sub> alkyl. Such purine or pyrimidine bases, analogues and derivatives are well known to those skilled in the art.

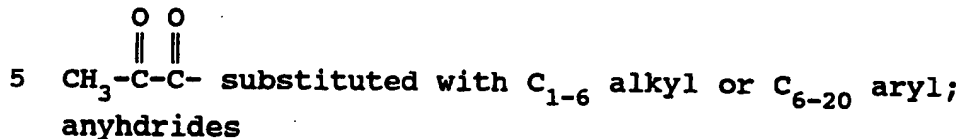
A "nucleoside analogue or derivative" is a 1,3-oxathiolane, 2,4-dioxolane or 1,3-dithiolane which has been modified in any of the following or combinations of the following ways: base modifications, such as addition of a substituent (e.g., 5-fluorocytosine) or replacement of one group by an isosteric group (e.g., 7-deazaadenine); sugar modifications, such as substitution of the C-2 and C-3 hydroxyl groups by any substituent, including hydrogen (e.g., 2',3'-dideoxynucleosides); alteration of the site of attachment of the sugar to the base (e.g., pyrimidine bases usually attached to the sugar at the N-1 site may be, for example, attached at the N-3 or C-6 site and purines usually attached at the N-9 site may be, for example, attached at N-7); alteration of the site of attachment of the base to the sugar (e.g., the base may be attached to the sugar at C-2, such as iso-DDA); or alteration of configuration of the sugar-base linkage (e.g., *cis* or *trans* configurations).

R<sub>3</sub> is a carbonyl substituted with hydrogen, hydroxyl, trialkylsilyl, trialkylsiloxy, C<sub>1-30</sub> alkyl, C<sub>7-30</sub> aralkyl, C<sub>1-30</sub> alkoxy, C<sub>1-30</sub> amine (primary,



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secondary or tertiary), C<sub>1-30</sub> thiol; C<sub>6-20</sub> aryl; C<sub>1-20</sub> alkenyl; C<sub>1-20</sub> alkynyl; 1,2-dicarbonyl, such as



such as  $\text{CH}_3-\overset{\text{O}}{\parallel}\text{C}-\text{O}-\overset{\text{O}}{\parallel}\text{C}-$  substituted with C<sub>1-6</sub> alkyl or C<sub>6-20</sub> aryl; azomethine substituted at nitrogen with hydrogen, C<sub>1-20</sub> alkyl or C<sub>1-10</sub> alkoxy or C<sub>1-10</sub> dialkylamino or at carbon with hydrogen, C<sub>1-20</sub> alkyl, or C<sub>1-20</sub> alkoxy; thiocarbonyl (C=S) substituted with hydroxyl, C<sub>1-20</sub> alkoxy, or C<sub>1-20</sub> thiol; a homologue of carbonyl, e.g.,



The preferred substituted carbonyl/carbonyl derivatives are alkoxy-carbonyls, such as methyl, ethyl, isopropyl, t-butyl and menthyl; carboxyls, diethylcarboxamide; pyrrolidine amide; methyl ketone and phenyl ketone. The more preferred substituted carbonyl/carbonyl derivatives are esters and carboxyls and the most preferred are esters.

R<sub>4</sub> is a chiral auxiliary. The term "chiral auxiliary" describes asymmetric molecules that are used to effect the chemical resolution of a racemic mixture. Such chiral auxiliaries may possess one chiral center such as methylbenzylamine or several chiral centers such as menthol. The purpose of the chiral auxiliary, once built into the starting material, is to allow simple separation of the resulting diastereomeric mixture. See, for example, J. Jacques et al.,

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Enantiomers, Racemates And Resolutions, pp. 251-369,  
John Wiley & Sons, New York (1981).

$R_5$ ,  $R_6$  and  $R_7$  are independently selected from the group consisting of hydrogen,  $C_{1-20}$  alkyl (e.g.,  
5 methyl, ethyl, t-butyl), optionally substituted by  
halogens (F, Cl, Br, I),  $C_{6-20}$  alkoxy (e.g., methoxy) or  
 $C_{6-20}$  aryloxy (e.g., phenoxy);  $C_{7-20}$  aralkyl (e.g.,  
benzyl), optionally substituted by halogen,  $C_{1-20}$  alkyl  
or  $C_{1-20}$  alkoxy (e.g., p-methoxybenzyl);  $C_{6-20}$  aryl  
10 (e.g., phenyl), optionally substituted by halogens,  
 $C_{1-20}$  alkyl or  $C_{1-20}$  alkoxy; trialkylsilyl; halogens (F,  
Cl, Br, I).

$R_8$  is selected from the group consisting of  
halogen (F, Cl, Br, I);  $C_{1-20}$  sulphonate esters,  
15 optionally substituted by halogens (e.g.,  
trifluoromethane sulphonate);  $C_{1-20}$  alkyl esters,  
optionally substituted by halogen (e.g.,  
trifluoroacetate); polyvalent halides (e.g.,  
triiodide); trisubstituted silyl groups of the general  
20 formula  $(R_5)(R_6)(R_7)Si$  (wherein  $R_5$ ,  $R_6$ , and  $R_7$  are as  
defined above); saturated or unsaturated selenenyl  
 $C_{6-20}$  aryl; substituted or unsubstituted  
 $C_{6-20}$  arylsulfenyl; substituted or unsubstituted  
 $C_{1-20}$  alkoxyalkyl; and trialkylsiloxo.

25  $L$  is a "leaving group", i.e., an atom or a  
group which is displaceable upon reaction with an  
appropriate purine or pyrimidine base, with or without  
the presence of a Lewis acid. Suitable leaving groups  
include acyloxy groups, alkoxy groups, e.g., alkoxy  
30 carbonyl groups such as ethoxy carbonyl; halogens such  
as iodine, bromine, chlorine, or fluorine; amido;  
azido; isocyanato; substituted or unsubstituted,  
saturated or unsaturated thiolates, such as thiomethyl  
or thiophenyl; substituted or unsubstituted, saturated

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or unsaturated seleno, seleninyl, or selenonyl compounds, such as phenyl selenide or alkyl selenide.

A suitable leaving group may also be -OR, where R is a substituted or unsubstituted, saturated or  
5 unsaturated alkyl group, e.g., C<sub>1-6</sub> alkyl or alkenyl group; a substituted or unsubstituted aliphatic or aromatic acyl group, e.g., a C<sub>1-6</sub> aliphatic acyl group such as acetyl and a substituted or unsubstituted aromatic acyl group such as benzoyl; a substituted or  
10 unsubstituted, saturated or unsaturated alkoxy or aryloxy carbonyl group, such as methyl carbonate and phenyl carbonate; substituted or unsubstituted sulphonyl imidazolidine; substituted or unsubstituted aliphatic or aromatic amino carbonyl group, such as  
15 phenyl carbamate; substituted or unsubstituted alkyl imide group such as trichloroacetamide; substituted or unsubstituted, saturated or unsaturated phosphonate, such as diethylphosphonate; substituted or unsubstituted aliphatic or aromatic sulphonyl or  
20 sulphonyl group, such as tosylate; or hydrogen.

As used in this application, the term "alkyl" represents a substituted (by a halogen, hydroxyl or C<sub>6-20</sub> aryl) or unsubstituted straight chain, branched chain, or cyclic hydrocarbon moiety having 1 to 30  
25 carbon atoms and preferably, from 1 to 6 carbon atoms.

The terms "alkenyl" and "alkynyl" represent substituted (by a halogen, hydroxyl or C<sub>6-20</sub> aryl) or unsubstituted straight, branched or cyclic hydrocarbon chains having 1 to 20 carbon atoms and preferably from  
30 1 to 5 carbon atoms and containing at least one unsaturated group (e.g., allyl).

The term "alkoxy" represents a substituted or unsubstituted alkyl group containing from 1 to 30 carbon atoms and preferably from 1 to 6 carbon atoms,  
35 wherein the alkyl group is covalently bonded to an

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adjacent element through an oxygen atom (e.g., methoxy and ethoxy).

The term "amine" represents alkyl, aryl, alkenyl, alkynyl, or aralkyl groups containing from 1 to 30 carbon atoms and preferably 1 to 12 carbon atoms, covalently bonded to an adjacent element through a nitrogen atom (e.g., pyrrolidine). They include primary, secondary and tertiary amines and quaternary ammonium salts.

The term "thiol" represents alkyl, aryl, aralkyl, alkenyl or alkynyl groups containing from 1 to 30 carbon atoms and preferably from 1 to 6 carbon atoms, covalently bonded to an adjacent element through a sulfur atom (e.g., thiomethyl).

The term "aryl" represents a carbocyclic moiety which may be substituted by at least one heteroatom (e.g., N, O, or S) and containing at least one benzenoid-type ring and preferably containing from 6 to 15 carbon atoms (e.g., phenyl and naphthyl).

The term "aralkyl" represents an aryl group attached to the adjacent atom by an alkyl (e.g., benzyl).

The term "alkoxyalkyl" represents an alkoxy group attached to the adjacent group by an alkyl group (e.g., methoxymethyl).

The term "aryloxy" represents a substituted (by a halogen, trifluoromethyl or C<sub>1-5</sub> alkoxy) or unsubstituted aryl moiety covalently bonded through an oxygen atom (e.g., phenoxy).

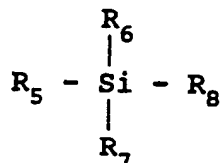
The term "acyl" refers to a radical derived from a carboxylic acid, substituted (by a halogen (F, Cl, Br, I), C<sub>6-20</sub> aryl or C<sub>1-6</sub> alkyl) or unsubstituted, by replacement of the -OH group. Like the acid to which it is related, an acyl radical may be aliphatic or aromatic, substituted (by a halogen, C<sub>1-5</sub>

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alkoxyalkyl, nitro or O<sub>2</sub>) or unsubstituted, and whatever the structure of the rest of the molecule may be, the properties of the functional group remain essentially the same (e.g., acetyl, propionyl, isobutanoyl, pivaloyl, hexanoyl, trifluoroacetyl, chloroacetyl, and cyclohexanoyl).

A key feature of the processes of this invention is the use of a substituted carbonyl or carbonyl derivative as R<sub>3</sub> instead of a protected hydroxymethyl group as previously described in the art. Surprisingly, the substituted carbonyl or carbonyl derivative is not cleaved by exposure to a Lewis acid, as would have been expected by one of skill in the art when a Lewis acid of formula (III) is added to a mixture of silylated purine or pyrimidine base and the chiral auxiliary-sugar compound obtained in Step 3. Instead, the substituted carbonyl/carbonyl derivative in the intermediate of formula (VI) forces the purine or pyrimidine base (R<sub>2</sub>) to add in the *cis*-configuration relative to the substituted carbonyl/carbonyl derivative group. Without a substituted carbonyl or carbonyl derivative attached to C4' (for example, when a hydroxymethyl group is instead used), the coupling procedures described in Step 4 will result in a mixture of *cis*- and *trans*-isomers.

Another key feature of the processes of this invention is the choice of Lewis acid. The Lewis acids used in the preparation of compounds of formula (I) have the general formula (III)



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wherein  $R_5$ ,  $R_6$ ,  $R_7$  and  $R_8$  are as defined previously. These Lewis acids may be generated in situ or prepared using any method known in the art (e.g., A.H. Schmidt, "Bromotrimethylsilane and Iodotrimethylsilane-Versatile  
5 Reagents for Organic Synthesis", Aldrichimica Acta, 14, pp. 31-38 (1981). The preferred Lewis acids of this invention are iodotrimethylsilane and trimethylsilyl triflate. The preferred  $R_5$ ,  $R_6$  and  $R_7$  groups are methyl or iodine. The most preferred  $R_5$ ,  $R_6$  and  $R_7$  group is  
10 methyl. The preferred  $R_8$  groups are iodine, chlorine, bromine or sulphonate esters. The most preferred  $R_8$  groups are iodine and trifluoromethane sulphonate.

In the preferred process of this invention, illustrated in Schemes 1 and 2, *cis*- and *trans*-isomers  
15 of a sugar of formula (II)



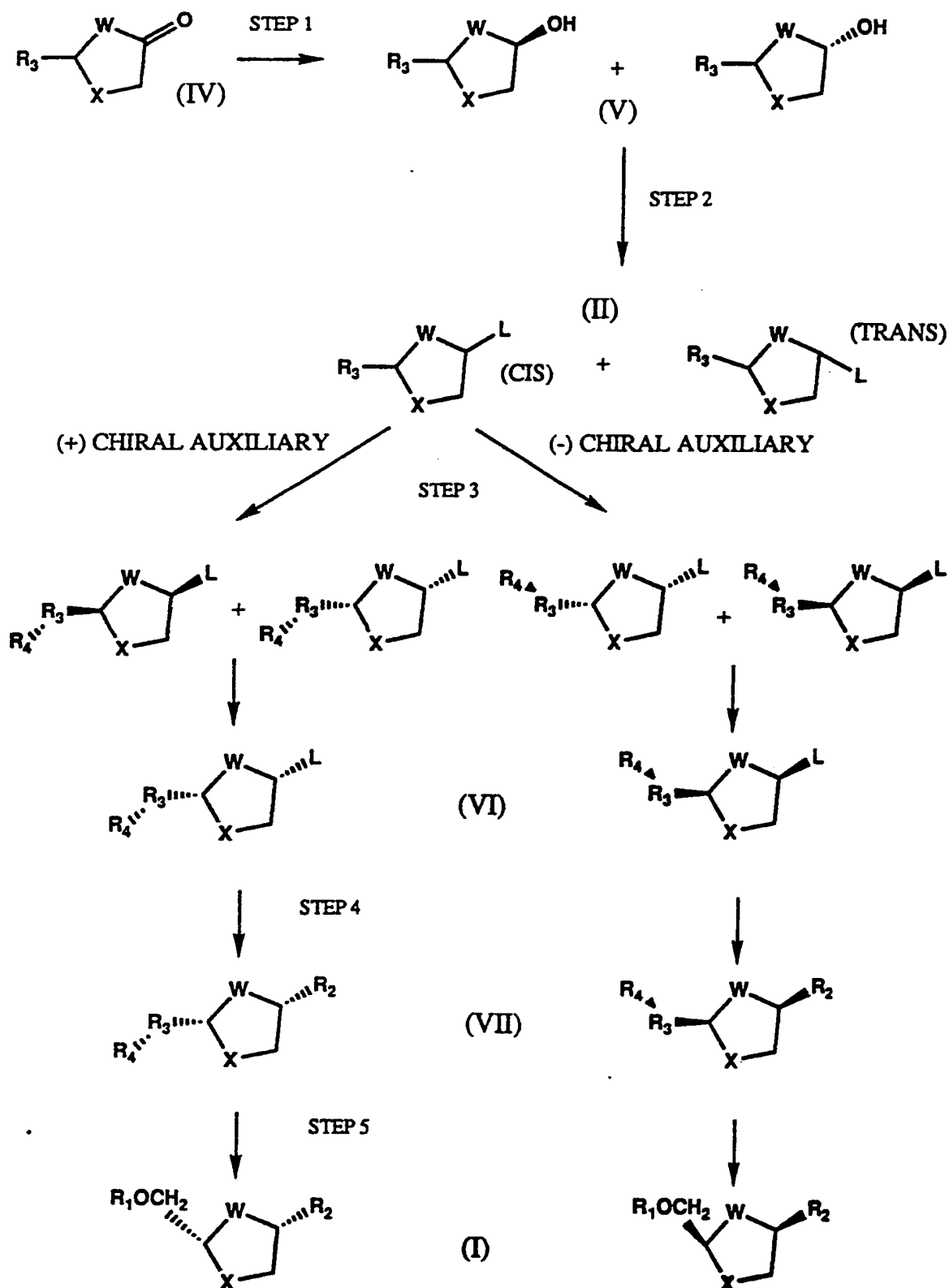
are separated by fractional crystallization and the desired configurational isomer selected. The selected *cis*- or the *trans*-isomer may then be resolved  
20 chemically, e.g., using a chiral auxiliary, enzymatically, or by other methods known in the art. The pure diastereomer is then coupled to a silylated purine or pyrimidine base in the presence of a Lewis acid to afford an optically active nucleoside of *cis*-  
25 configuration which is subsequently reduced to give a nucleoside of formula (I).

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Schemes 1A and 1B depict this preferred process as applied to any 1,3-oxathiolane, 2,4-dioxolane or 1,3-dithiolane.

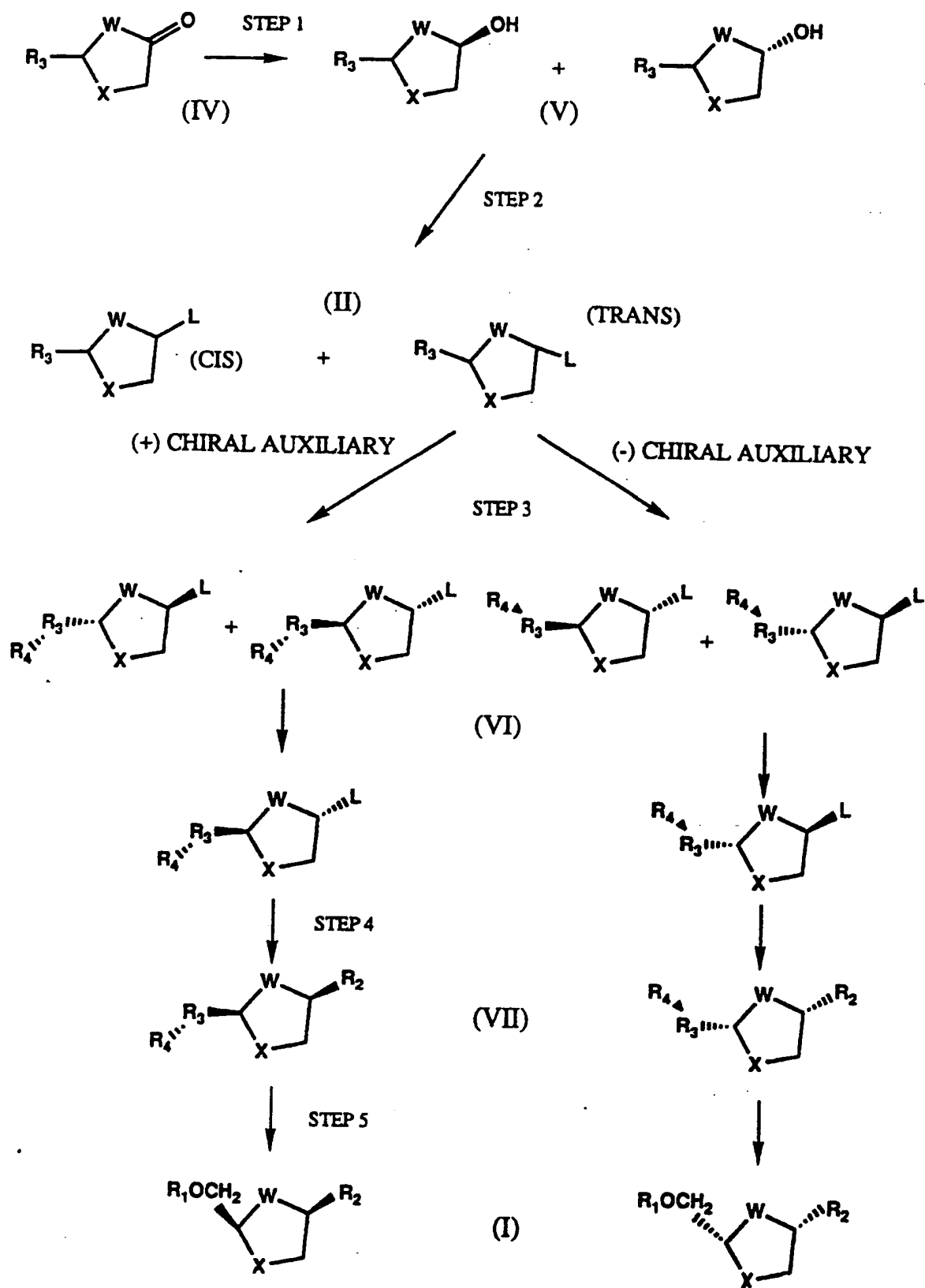
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## SCHEME 1A





## SCHEME 1B



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The various steps as illustrated in Schemes 1A and 1B may be briefly described as follows:

Step 1: The starting carbonyl-sugar of formula (IV) can be prepared by any method known in the art. E.g., J.M. McIntosh et al., "2-Mercaptoaldehyde Dimers and 2,5-Dihydrothiophenes from 1,3-oxathiolan-5-ones", Can. J. Chem., 61, pp. 1872-1875 (1983). The carbonyl group of this starting compound is reduced chemoselectively with a suitable reducing agent, such as disiamylborane to give the *cis*- and *trans*-isomers of formula (V). Ordinarily, less *cis*-isomer is produced than *trans*.

Step 2: The hydroxyl group in the intermediate of formula (V) is readily converted to a leaving group by any method known in the art (e.g., T.W. Greene Protective Groups In Organic Synthesis, pp. 50-72, John Wiley & Sons, New York (1981)) to give the novel intermediates of formula (II).

This anomeric mixture is then separated by fractional crystallization into the two configurational isomers. The solvent may be adjusted to select for either the *cis*- or *trans*-isomer. D.J. Pasto and C.R. Johnson, Organic Structure Determination, pp. 7-10, Prentice-Hall, Inc., New Jersey (1969).

Step 3: Either the *cis*- (Scheme 1A) or *trans*-isomer (Scheme 1B) of formula (II) is chemically resolved using a chiral auxiliary ( $R_4$ ). A suitable chiral auxiliary is one of high optical purity and where the mirror image is readily available, such as d- and l-menthol. The resulting diastereomers of formula (VI) are easily separated by fractional crystallization. Alternatively, either the *cis*- or the *trans*-isomer may be resolved enzymatically or by other methods known in the art. Jacques et al., Enantiomers,

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Racemates And Resolutions, pp. 251-369, John Wiley & Sons, New York (1981).

The optical purity of the diastereomer (VI, VII or I) can be determined by chiral HPLC methods, specific rotation measurements and NMR techniques. As a general rule, if the opposite enantiomer is desired, it may be obtained by using the mirror image of the chiral auxiliary initially employed. For example, if the chiral auxiliary d-menthol produces a (+)-enantiomer nucleoside, its mirror image, l-menthol, will produce the (-)-enantiomer.

Step 4: A previously silylated (or silylated in situ) purine or pyrimidine base or analogue or derivative thereof is then glycosylated with the resulting pure diastereomer in the presence of a Lewis acid of formula (III), such as iodotrimethylsilane (TMSI) or trimethylsilyl triflate (TMSOTf), to give a nucleoside of *cis*-configuration of formula (VII). This nucleoside is optically active and is substantially free of the corresponding *trans*-isomer (i.e., it contains less than 20%, preferably no more than 10% and more preferably no more than 5% of the *trans*-isomer).

The preferred silylating agent for pyrimidine bases are t-butyldimethylsilyl triflate 1,1,1,3,3,3 hexamethyldisilazane and trimethylsilyl triflate. It is believed that the bulky t-butyl group increases yields by weakening the interaction between the Lewis acid and silylated pyrimidine base.

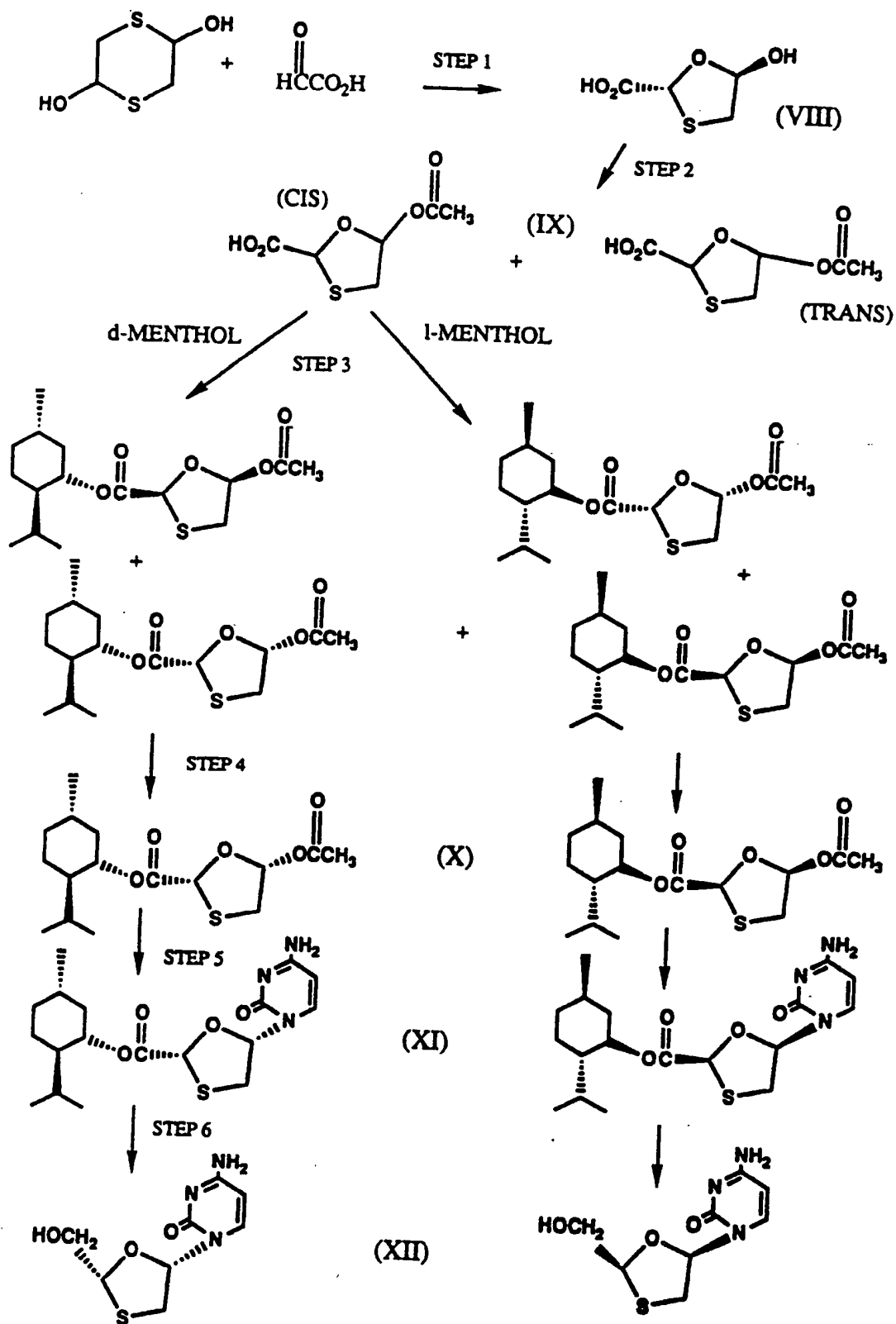
The preferred method of mixing reagents in Step 4 is to first add the chiral auxiliary-sugar of formula (VI) to the silylated purine or pyrimidine base. The Lewis acid of formula (III) is then added to the mixture.

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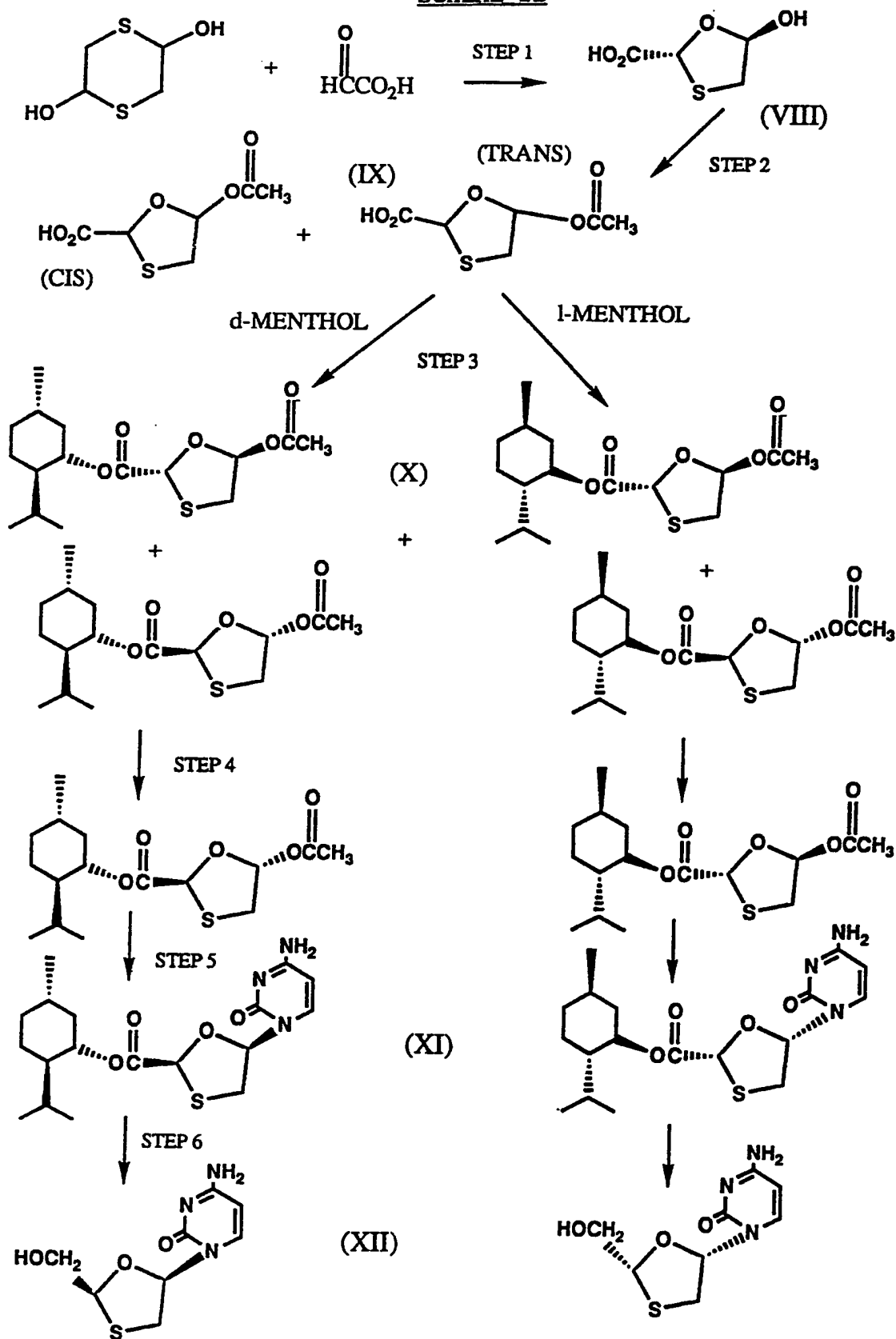
Step 5: The *cis*-nucleoside obtained in Step 4 may then be reduced with an appropriate reducing agent to remove the chiral auxiliary and give a specific stereoisomer of formula (I). The absolute configuration of this stereoisomer corresponds to that of the nucleoside intermediate of formula (VII). As shown in Scheme 1, either the *cis*- (Scheme 1A) or the *trans*-isomers (Scheme 1B) obtained in Step 2 will yield a *cis* end product.

10 Schemes 2A and 2B illustrate the application of the process of Schemes 1A and 1B to the synthesis of the enantiomers of *cis*-2-hydroxymethyl-5-(cytosin-1'-yl)-1,3-oxathiolanes. Although this process is illustrated using specific reagents and starting materials, it will be appreciated by one of skill in the art that suitable analogous reactants and starting materials may be used to prepare analogous compounds.

## SCHEME 2A



## SCHEME 2B



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The various steps illustrated in Schemes 2A and 2B may be briefly described as follows:

Step 1: A mercaptoacetaldehyde monomer, preferably produced from a dimer, such as 2,5-dihydroxy-1,4-dithiane, in an appropriate solvent (preferably *t*-butylmethyl ether) is reacted with glyoxylic acid to give exclusively the *trans*-hydroxy acid of formula (VIII).

Step 2: The acid of formula (VIII) is reacted with an acid chloride, such as acetyl chloride in the presence of pyridine and an acylation catalyst, such as 4-dimethylaminopyridine, or preferably with an acid anhydride such as acetic anhydride in the presence of acetic acid and an acylation catalyst, such as sulfuric acid, to give a diastereomeric mixture of *cis*- and *trans*-acetoxy acids of formula (IX).

The racemic diastereomeric acid mixture obtained in Step 2 is fractionally crystallized using any combination of solvents (preferably benzene and ether) to give exclusively either the *cis*- or the *trans*-acetoxy acid of formula (IX) each as a racemic mixture.

Step 3: Either the *cis*- or the *trans*-acetoxy acid of formula (IX) is reacted with an appropriate chiral auxiliary preferably, 1-menthol or *d*-menthol, in a suitable organic solvent, such as dichloromethane, using an activating agent, such as dicyclohexylcarbodiimide, and an esterification catalyst, such as 4-dimethylaminopyridine, to give a diastereomeric mixture of the *cis*- or *trans*-esters respectively.

Alternatively, the compound of formula (IX) may be converted to an acid chloride by any means known in the art, such as with oxalyl chloride in an appropriate solvent, e.g., dichloromethane or *N,N*-dimethylformamide. The acid chloride is then reacted

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with a chiral auxiliary in a suitable organic solvent using an esterification catalyst.

Step 4: The above diastereomeric mixture of either the *cis*- or the *trans*-esters is fractionally  
5 crystallized using any combination of solvents (preferably ether and petroleum ether (40-60°C)) preferably at low temperature to give exclusively the *cis*- or the *trans*-acetoxy menthyl ester of formula (X), respectively.

10 Step 5: Either the *cis*- or the *trans*-acetoxy compound of formula (X) is reacted with cytosine or other purine or pyrimidine base or analogue thereof. The purine or pyrimidine base or analogue is preferably previously silylated with hexamethyldisilazane or more  
15 preferably silylated in situ with *t*-butyldimethylsilyl triflate in a compatible organic solvent, such as dichloromethane containing a hindered base preferably 2,4,6-collidine. A Lewis acid of formula (III), preferably iodotrimethylsilane or trimethylsilyl  
20 triflate, is then added to give the *cis*-compound of formula (XI) in a highly diastereoselective manner.

Step 6: The optically active *cis*-nucleoside of formula (XI) is reduced stereospecifically with a reducing agent preferably lithium triethylborohydride  
25 or more preferably lithium aluminum hydride in an appropriate solvent such as tetrahydrofuran or diethyl ether to give the compound of formula (XII) and menthol.

A second process for the diastereoselective  
30 synthesis of compounds of formula (I) is illustrated by Schemes 3A and 3B and 4A and 4B. In the process of Schemes 3A and 3B carbonyl-sugar with an  $R_3$  substituent at C4' is reacted with a chiral auxiliary ( $R_4$ ) to give a diastereomeric mixture of two optically active chiral



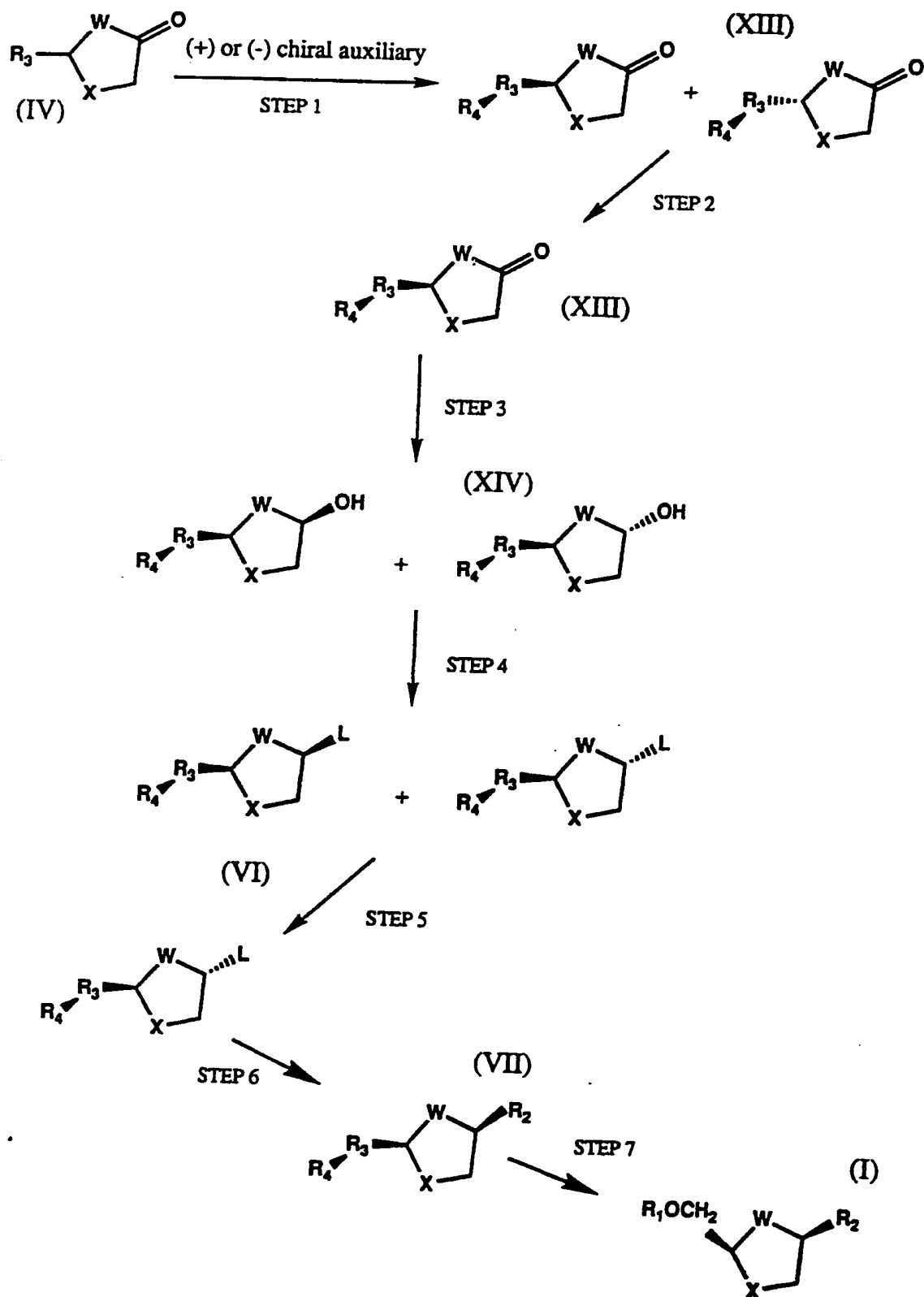
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auxiliary sugars. The actual diastereomer produced depends on whether the (+) or (-) chiral auxiliary is used. This optically active mixture may be chemoselectively reduced and the resulting hydroxyl  
5 group converted to a leaving group to afford a diastereomeric mixture of four chiral auxiliary-sugars, two in the *cis*- configuration and two in the *trans*- configuration (Scheme 3B). Subsequent fractional crystallization gives a single diastereomer.

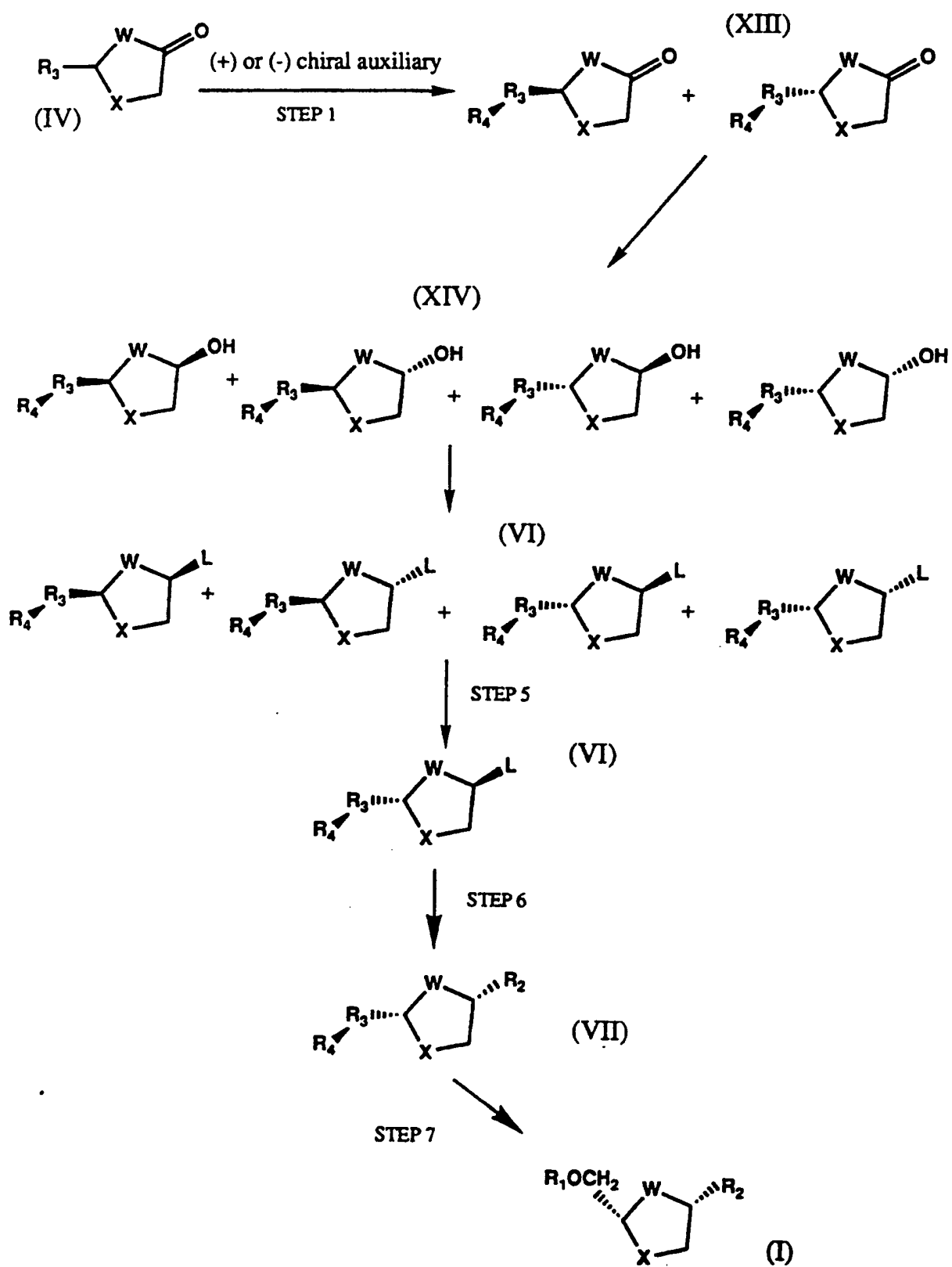
10 Alternatively, the optically active mixture of chiral auxiliary-sugars may first be separated by chromatography or fractional crystallization and then reduced and the resulting hydroxyl group converted to a leaving group (Scheme 3A). Subsequent fractional  
15 crystallization yields any desired diastereomer. The solvent may be adjusted to select for either the *cis*- or the *trans*-isomer. Each isolated optically active diastereomer may be carried on further to compounds of formula (I) in a manner analogous to that described in  
20 Schemes 1 and 2.

Schemes 3A and 3B depict the second process of this invention as applied to any 1,3-oxathiolane, 2,4-dioxolane or 1,3-dithiolane.

## SCHEME 3A



## SCHEME 3B



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The various steps involved in the synthesis of the nucleosides of formula (I) as depicted in Schemes 3A may be briefly described as follows:

Step 1: The starting material of formula (IV), prepared by any method known in the art, is reacted with a chiral auxiliary (see, e.g., T.W. Greene, "Protective Groups in Organic Synthesis", John Wiley and Sons, New York (1981) to yield a mixture of two diastereomers of formula (XIII). The particular mixture produced will depend on which chiral auxiliary (+ or -) is used.

Step 2: The mixture of two diastereomers of formula (XIII) is separated by fractional crystallization or chromatography to yield one diastereomer of formula (XIII).

Step 3: The single isomer of formula (XIII) is chemoselectively reduced by a suitable reducing agent, such as disiamylborane to give a mixture of two diastereomers of formula (XIV).

Step 4: The hydroxyl groups of the two diastereomers of formula (XIV) are converted to leaving groups by any method known in the art to give a mixture of two diastereomers of formula (VI).

Step 5: Either the *cis*- or *trans*-isomer is separated out of the mixture of two diastereomers of formula (VI), as obtained in Step 4, by fractional crystallization or chromatography. The solvent may be adjusted to select for the *cis*- or *trans*-isomer.

Step 6: The single diastereomer of formula (VI) is reacted with previously silylated (or silylated in situ) purine or pyrimidine base or analogue or derivative. Then, addition of a Lewis acid of formula (III), such as iodotrimethylsilane (TMSI) or trimethylsilyl triflate (TMSOTf) yields a nucleoside of

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*cis*-configuration of formula (VII). This nucleoside is substantially free of the corresponding *trans*-isomer.

Step 7: The optically active *cis*-nucleoside of formula (VII) is reduced stereospecifically with a  
5 reducing agent preferably lithium triethylborohydride or more preferably lithium aluminum hydride in an appropriate solvent such as tetrahydrofuran or diethyl ether to give the compound of formula (I) and menthol.

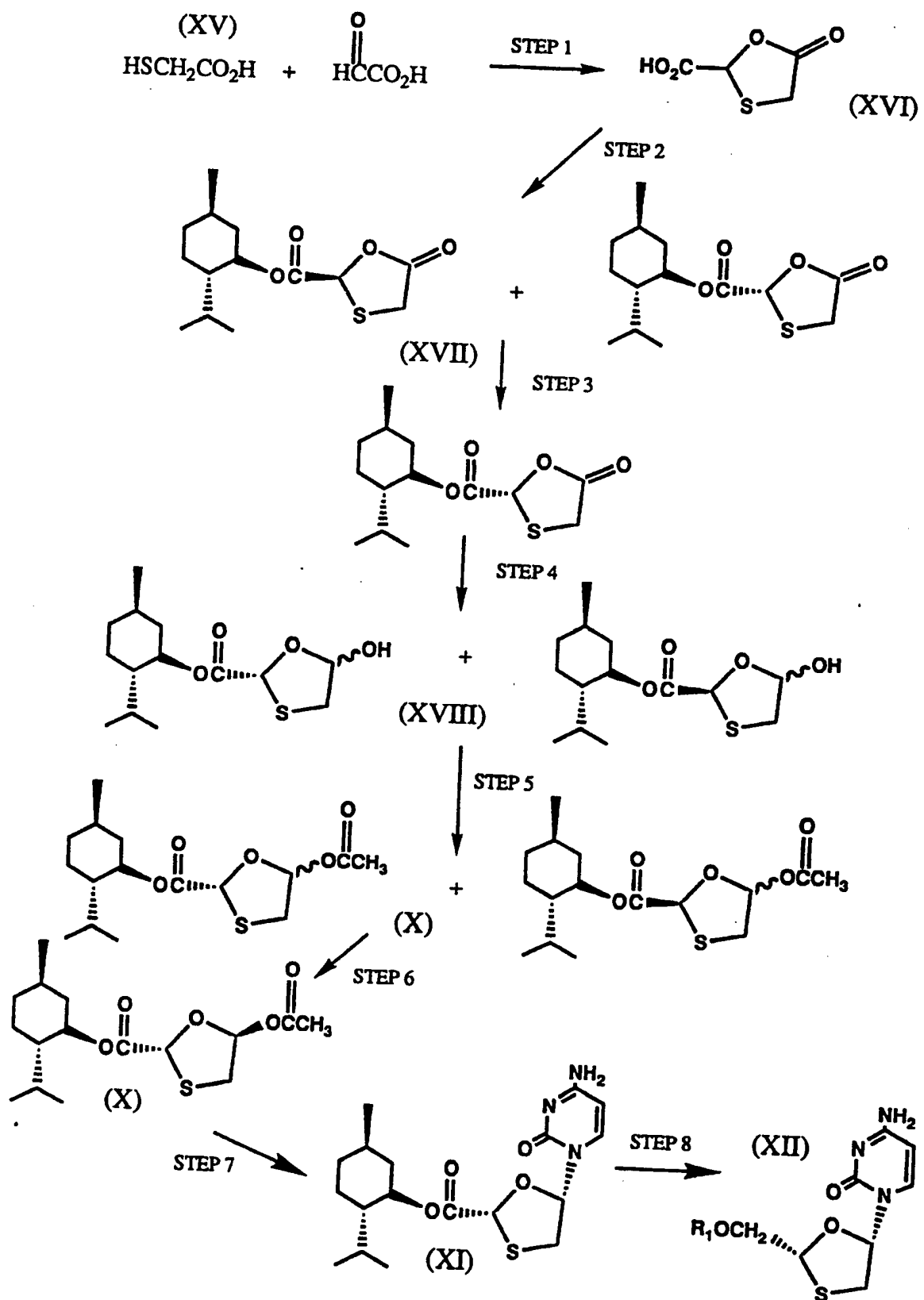
Alternatively, as shown in Scheme 3B, the  
10 mixture of diastereomers of formula (XIII) is chemoselectively reduced with a suitable reducing agent, such as disiamylborane to give a mixture of four diastereomers of formula (XIV). The hydroxyl groups in this mixture of four diastereomers of formula (XIV) are  
15 converted to leaving groups any method in the art to afford a mixture of four diastereomers of formula (VI). Either a *cis*- or a *trans*-isomer of formula (VI) is separated out of the mixture of four diastereomers of formula (VI) by fractional crystallization or  
20 chromatography. The solvent may be adjusted to select for a *cis*- or *trans*-isomer. The single diastereomer of formula (VI) is reacted with previously silylated (or silylated in situ) purine or pyrimidine base or analogue or derivative. Then, addition of a Lewis acid  
25 of formula (III), such as iodotrimethylsilane (TMSI) or trimethylsilyl triflate (TMSOTf) affords a nucleoside of *cis*- configuration of formula (VII) which is reduced with an appropriate reducing agent to give a specific stereoisomer of formula (I).

30 Schemes 4A and 4B illustrate the application of the process of Scheme 3 to the synthesis of the enantiomers of *cis*-2-hydroxymethyl-5-(cytosin-1'-yl)-1,3-oxathiolanes. Although this process is illustrated using specific reagents and starting materials, it will

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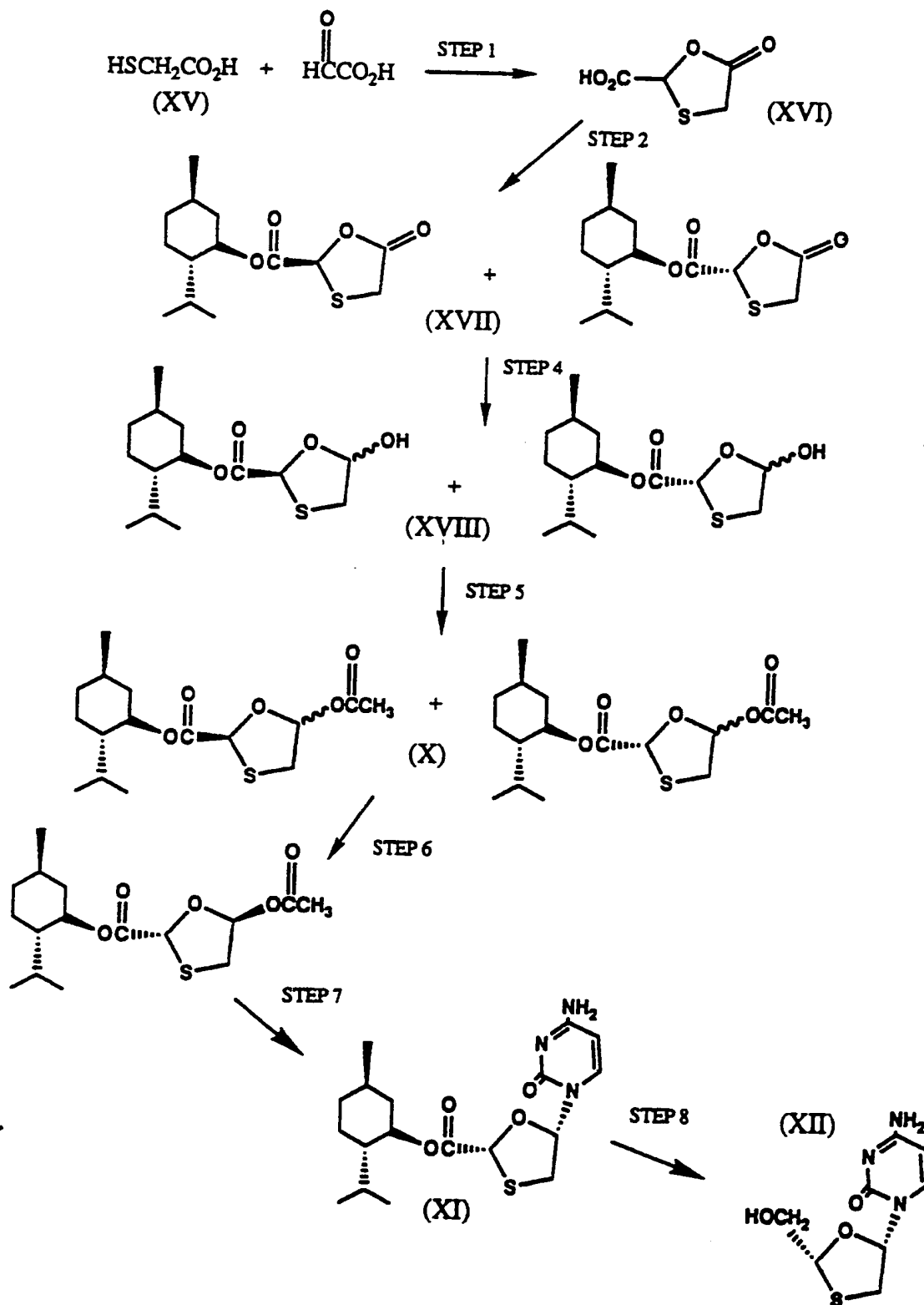
be appreciated by one of skill in the art that suitable analogous reactants and starting materials may be used to prepare analogous compounds.

## SCHEME 4A



- 30 -

## SCHEME 4B





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The various steps involved in the synthesis of the nucleosides of formula (I) as depicted in Scheme 4 may be briefly described as follows:

Step 1: The known mercaptoacetic acid of formula (XV) is reacted with an appropriate aldehyde of formula  $R_3CHO$ , wherein  $R_3$  is preferably an alkoxy carbonyl, such as menthyl glyoxylate and more preferably a carboxyl group, such as glyoxylic acid (see e.g., J.M. McIntosh et al., "2-Mercaptoaldehyde Dimers and 2,5-Dihydrothiophenes from 1,3-oxathiolan-5-ones", Can. J. Chem., 61, pp. 1872-1875 (1983)) in a compatible organic solvent, such as toluene, to give the intermediate of formula (XVI).

Step 2: The compound of formula (XVI) is reacted with an appropriate chiral auxiliary, preferably l-menthol or d-menthol in a compatible organic solvent, such as dichloromethane, using an activating agent, such as dicyclohexylcarbodiimide, and an esterification catalyst, such as 4-dimethylamino-pyridine, to give the compounds of formula (XVII).

Step 3: The diastereomeric compounds of formula (XVII) are preferably separated by fractional crystallization (Scheme 4A), but may be carried on further without separation (Scheme 4B).

Step 4: The compounds of formula (XVII) are reduced with an appropriate reducing agent such as disiamylborane in a compatible organic solvent, such as tetrahydrofuran (A. Pelter et al., "Borane Reagents", Academic Press, p. 426 (1988)), to give the compounds of formula (XVIII).

Step 5: The compounds of formula (XVIII) are reacted with an acid chloride or acid anhydride, such as acetic anhydride, in the presence of pyridine and an acylation catalyst, such as 4-dimethylaminopyridine, to give the compounds of formula (X).

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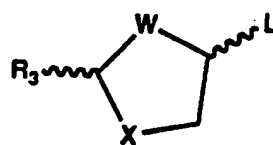
Step 6: The diastereomeric compounds of formula (X), if not already separated (Scheme 4A), are now separated preferably by fractional crystallization (Scheme 4B) to give either the *cis*- or the *trans*-  
5    acetoxy compound of formula (X).

Step 7: Either the *cis*- or the *trans*-acetoxy compound of formula (X) is reacted with cytosine or other purine or pyrimidine base or analogue thereof. The purine or pyrimidine base or analogue is  
10    preferably previously silylated with hexamethyldisilazane or more preferably silylated in situ with *t*-butyldimethylsilyl triflate in a compatible organic solvent, such as dichloromethane containing a hindered base preferably 2,4,6-collidine. A Lewis acid,  
15    preferably one derived from the compounds of formula (III), more preferably iodotrimethylsilane or trimethylsilyl triflate, is then added to give the *cis* compound of formula (XI) in a highly diastereoselective manner.

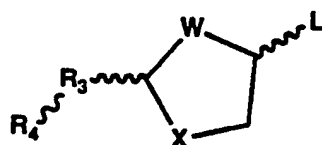
20    Step 8: The optically active *cis*-nucleoside of formula (XI) is reduced stereospecifically with a reducing agent, preferably lithium triethylborohydride, or more preferably, lithium aluminum hydride, in an appropriate solvent, such as tetrahydrofuran or diethyl  
25    ether, to give the compound of formula (XII).

In the diastereoselective processes of this invention, the following intermediates are of particular importance:

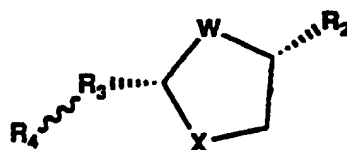
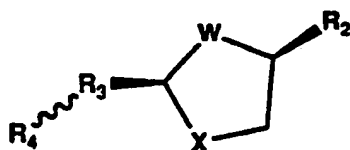
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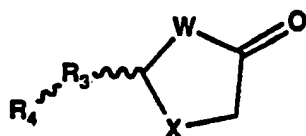
(II)



(VI)

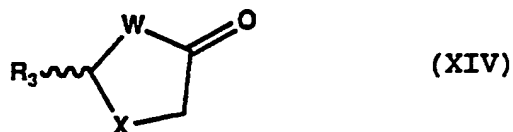


(VII)



(XIII)

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wherein  $R_3$ ,  $R_4$  and L are as defined above;

*trans*-5-hydroxyoxathiolane-2-carboxylic acid;

(1'R,2'S,5'R)-menthyl-1,3-oxathiolan-5-one-

5 2S-carboxylate;

(1'R,2'S,5'R)-menthyl-1,3-oxathiolan-5-one-

2R-carboxylate;

(1'R,2'S,5'R)-menthyl-5S-hydroxy-1,3-

oxathiolane-2S-carboxylate;

10 (1'R,2'S,5'R)-menthyl-5R-hydroxy-1,3-

oxathiolane-2R-carboxylate;

(1'R,2'S,5'R)-menthyl-5S-hydroxy-1,3-

oxathiolane-2R-carboxylate;

(1'R,2'S,5'R)-menthyl-5R-hydroxy-1,3-

15 oxathiolane-2S-carboxylate;

(1'R,2'S,5'R)-menthyl-5S-acetoxy-1,3-

oxathiolane-2S-carboxylate;

(1'R,2'S,5'R)-menthyl-5R-acetoxy-1,3-

oxathiolane-2R-carboxylate;

20 (1'R,2'S,5'R)-menthyl-5S-acetoxy-1,3-

oxathiolane-2R-carboxylate;

(1'R,2'S,5'R)-menthyl-5R-acetoxy-1,3-

oxathiolane-2S-carboxylate;

(1'S,2'R,5'S)-menthyl-5R-acetoxy-1,3-

25 oxathiolane-2S-carboxylate;

(1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-

oxathiolane-2R-carboxylate;

(1'S,2'R,5'S)-menthyl-5R-acetoxy-1,3-

oxathiolane-2R-carboxylate;

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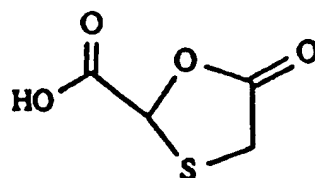
- (1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolane-2S-carboxylate;
- (1'R,2'S,5'R)-menthyl-5S-(cytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate;
- 5 (1'S,2'R,5'S)-menthyl-5S-(cytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate;
- (1'R,2'S,5'R)-menthyl-5R-(cytosin-1"-yl)-1,3-oxathiolane-2S-carboxylate;
- (1'S,2'R,5'S)-menthyl-5R-(cytosin-1"-yl)-1,3-oxathiolane-2S-carboxylate;
- 10 (1'R,2'S,5'R)-menthyl-5R-(5"-fluorocytosin-1"-yl)-1,3-oxathiolane-2S-carboxylate;
- (1'S,2'R,5'S)-menthyl-5S-(5"-fluorocytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate;
- 15 (1'S,2'R,5'S)-menthyl-5S-(N-4"-acetylcytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate;
- (1'R,2'S,5'R)-menthyl-5S-(cytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate;
- (1'S,2'R,5'S)-menthyl-1,3-oxathiolane-2R-carboxylate;
- 20 (1'S,2'R,5'S)-menthyl-4R-hydroxy-1,3-oxathiolane-2R-carboxylate and (1'S,2'R,5'S)-menthyl-4S-hydroxy-1,3-oxathiolane-2R-carboxylate;
- (1'S,2'R,5'S)-menthyl-4R-chloro-1,3-oxathiolane-2R-carboxylate and (1'S,2'R,5'S)-menthyl-4S-chloro-1,3-oxathiolane-2R-carboxylate;
- 25 *cis*-2(N-methyl-N-methoxyaminocarbonyl)-5-(uracil-1'-yl)-1,3-oxathiolane;
- cis*- and *trans*-2-benzoyl-5-acetoxy-1,3-oxathiolane;
- 30 *cis*-2-(1'-pyrrolidinocarbonyl)-5-acetoxy-1,3-oxathiolane;
- cis*-2-carbomethoxy-5-(5'-bromouracil-1'-yl)-1,3-oxathiolane;

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- cis*-2-carboxyl-5-(uracil-1'-yl)-1,3-oxathiolane;  
*cis*-2-(1'-pyrrolidinocarbonyl)-5-(uracil-1'-yl)-1,3-oxathiolane;  
5 *cis* 2-benzoyl-5-(uracil-1'-yl)-1,3-oxathiolane;  
*cis*- and *trans*-isopropyl 5-acetoxy-1,3-oxathiolane-2-carboxylate;  
*cis*-isopropyl-5-(cytosin-1'-yl)-1,3-oxathiolane-2-carboxylate;  
10 *cis*- and *trans*-*t*-butyl 5-acetoxy-1,3-oxathiolane-2-carboxylate;  
*cis*-*t*-butyl-5-(cytosin-1'-yl)-1,3-oxathiolane-2-carboxylate;  
15 *cis*- and *trans*-2-*N,N*-diethylaminocarbonyl- 5-acetoxy-1,3-oxathiolane;  
*cis*-2-*N,N*-diethylaminocarbonyl-5-(cytosin-1'-yl)-1,3 -oxathiolane;  
*cis*- and *trans*-2-carboethoxy-4-acetoxy-1,3-oxathiolane;  
20 dioxolane;  
*cis*- and *trans*-2-carboethoxy-4-(thymine-1'-yl)-1,3-dioxolane; and  
*cis*- and *trans*-2-carboethoxy-4-(*N*-4'-acetylcytosine-1'-yl)-1,3-dioxolane.

- 25 The following examples illustrate the present invention in a manner of which it can be practiced but, as such, should not be construed as limitations upon the overall scope of the processes of this invention. Except where specifically noted, all  $[\alpha]_D$  measurements  
30 were recorded at ambient temperature.

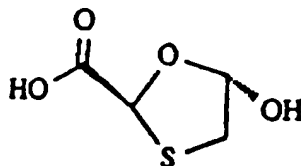
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EXAMPLE 11,3-OXATHIOLAN-5-ONE-2-CARBOXYLIC ACID

(XVI)

Toluene (700 mL), mercaptoacetic acid (38 mL,  
5 50.03 g, 0.543 mol), and p-toluenesulfonic acid (1.0 g)  
were added to a solution of glyoxylic acid monohydrate  
(50.0 g, 0.543 mol) in 200 mL of THF in a 2 L round  
bottom flask equipped with a Dean-Stark trap and  
condenser. The resultant reaction mixture was refluxed  
10 for 3 hours until 24.0 mL of H<sub>2</sub>O was azeotropically  
removed. The reaction mixture was cooled, followed by  
removal of solvent under reduced pressure to yield an  
off-white solid. This material was purified by  
recrystallization (hexanes-EtOAc) to give 60.0 g of the  
15 product as a crystalline white solid: m.p. 140-143°C;  
<sup>1</sup>H NMR (DMSO) δ 3.84 (q, 2H, JAB=16.7 Hz), 6.00 (s,  
1H).

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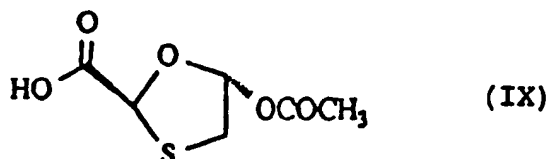
Example 2TRANS-5-HYDROXYOXATHIOLANE-2-CARBOXYLIC ACID

(VIII)

A suspension of dithian-1,4-diol (82.70 g, 0.54 mol) and glyoxylic acid monohydrate (100.0 g, 1.09 mol) in tert-butyl methyl ether (1.1 L) was stirred under a blanket of nitrogen and heated to reflux under Dean and Stark conditions. The reflux was continued for 8 hours during which time 15.3 mL (0.85 mol) of water was collected. The slightly turbid mixture was filtered, and the solvent was distilled at atmospheric pressure until a volume of 600 mL remained. Cyclohexane (340 mL) was added and the solution was cooled to 5°C, seeded, and allowed to stir and crystallize. The suspension was stirred at 0-5°C for 2 hours. The product was isolated by filtration, washed with 100 mL of tert-butyl methyl ether-cyclohexane (2:1), and was dried overnight in vacuo at room temperature (94.44 g): m.p. 94.5°C; <sup>1</sup>H NMR (DMSO)  $\delta$  2.85 (dd, 1H, J=2.4, 10.5 Hz), 3.13 (dd, 1H, J=4.3, 10.5 Hz), 5.47 (s, 1H), 5.84 (brs, 1H), 6.95 (d, 1H, J=4.7 Hz).

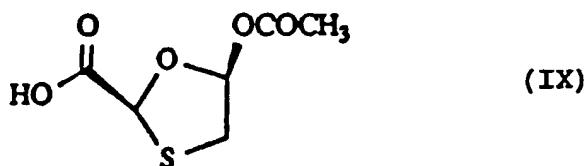


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Example 3TRANS-5-ACETOXY-1,3-OXATHIOLANE-2-CARBOXYLIC-ACID

One drop of concentrated  $\text{H}_2\text{SO}_4$  was added to a  
5 thoroughly stirred solution of trans-5-hydroxy-  
oxathiolane-2-carboxylic acid (7.0 g, 46.7 mmol) in  
glacial acetic acid (40 mL) and acetic anhydride  
(15 mL, 15.9 mmol) at ambient temperature. The  
resultant clear solution was stirred for 1 hour and  
10 then poured onto crushed ice and brine (20 mL). This  
mixture was extracted with  $\text{CH}_2\text{Cl}_2$  (100 mL) and the  
combined extract was dried over anhydrous magnesium  
sulfate. The solvent was removed under reduced  
pressure to give 8.5 g (95%) of a light yellow syrup  
15 which consisted of trans- and cis-5-acetoxy-1,3-  
oxathiolane-2-carboxylic acid in a 2:1 ratio. The  
mixture was dissolved in benzene (20 mL) and was left  
standing overnight during which white crystals were  
formed. A small amount of ether was added and the  
20 solid was collected by filtration and washed with more  
ether to give 2 g (22%) of trans-5-acetoxy-1,3-  
oxathiolane-2-carboxylic acid: m.p.  $111.3^\circ\text{C}$ ;  $^1\text{H}$  NMR  
(DMSO)  $\delta$  2.03 (s, 3H), 3.21 (d, 1H,  $J=12$  Hz), 3.32 (dd,  
1H,  $J=3, 12$  Hz), 5.65 (s, 1H), 6.65 (d, 1H,  $J=4$  Hz);  $^{13}\text{C}$   
25 NMR (DMSO)  $\delta$  20.91, 36.51, 78.86, 99.15, 169.36,  
170.04.

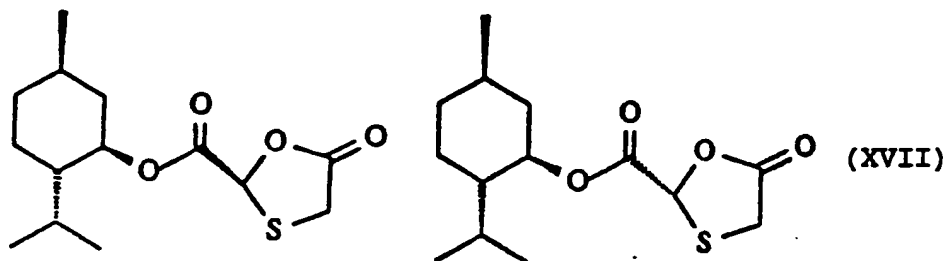
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Example 4CIS-5-ACETOXY-1,3-OXATHIOLANE-2-CARBOXYLIC ACID

The filtrate obtained from Example 3 was concentrated under reduced pressure and redissolved in ether. This solution was kept at room temperature and *cis*-5-acetoxy-1,3-oxathiolane-2-carboxylic acid slowly crystallized out as a white solid (2.1 g, 23%): m.p. 111.7°C; <sup>1</sup>H NMR (DMSO) δ 1.96 (s, 3H), 3.25-3.33 (m, 2H), 5.74 (s, 1H), 6.69 (d, 1H, J=3 Hz); <sup>13</sup>C NMR (DMSO) δ 21.0, 37.16, 79.57, 98.58, 169.36, 170.69.

Example 5

(1'R,2'S,5'R)-MENTHYL-1,3-OXATHIOLAN-5-ONE-2S-CARBOXYLATE AND (1'R,2'S,5'R)-MENTHYL-1,3-OXATHIOLAN-5-ONE-2R-CARBOXYLATE



Oxalyl chloride (11 mL, 123.6 mmol) was added through a dropping funnel over a period of 30 minutes to a stirred solution of 1,3-oxathiolan-5-one-2-

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carboxylic acid (12.2 g, 82.4 mmol) in anhydrous THF (20 ml) and  $\text{CH}_2\text{Cl}_2$  (40 mL) at room temperature under an argon atmosphere. The resultant solution was heated at 65°C for 30 minutes and then was concentrated in vacuo to give an oily product (11.6 g, 90%). The crude acid chloride obtained was redissolved in dry  $\text{CH}_2\text{Cl}_2$  (40 mL) and cooled at 0°C. (1R,2S,5R)-menthol (12.8 g, 82.4 mmol) dissolved in  $\text{CH}_2\text{Cl}_2$  (25 mL) was slowly added to this cooled solution. The resultant solution was stirred at room temperature overnight. The reaction mixture was diluted with  $\text{CH}_2\text{Cl}_2$  (200 mL) and washed with water, saturated aqueous  $\text{NaHCO}_3$  solution, brine, and then was dried over anhydrous  $\text{Na}_2\text{SO}_4$ . The solvent was removed and the crude product thus obtained was filtered through a short silica column (100 g, Merck) eluted with EtOAc-hexanes. Concentration of the appropriate fractions gave a 1:1 mixture of (1'R,2'S,5'R)-menthyl-1,3-oxathiolan-5-one-2S-carboxylate and (1'R,2'S,5'R)-menthyl-1,3-oxathiolan-5-one-2R-carboxylate (20 g, 84.7% overall) as a viscous oil:  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  0.77 (3H), 0.91 (6H), 1.00-1.15 (2H), 1.40-2.10 (6H), 3.56 (1H), 3.82 (1H), 4.80 (1H). 5.62 (1H);  $^{13}\text{C}$  NMR  $\delta$  16.7, 21.2, 21.3, 22.5, 23.80, 23.84, 26.7, 26.8, 30.6, 31.91, 31.94, 34.57, 40.6, 41.07, 47.5, 47.6, 74.1, 74.2, 77.7, 168.1, 172.8.

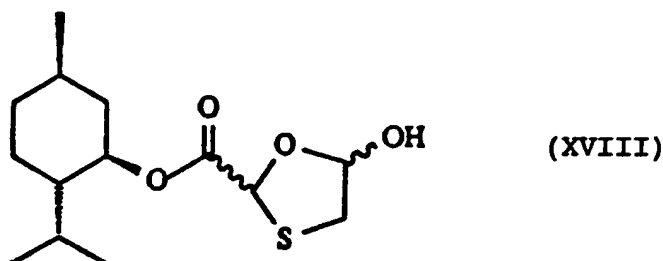
The above mixture (20 g) was dissolved in a minimum amount of pentane-petroleum ether (40-60°C) (1:2, 30 mL). The resultant solution was cooled at -70°C for 10 minutes and the crystalline compound that was formed was quickly collected by filtration and washed with more cold petroleum ether (10 mL). This crystalline compound, isolated in 12.5% yield, was found to consist of one isomer as indicated by  $^1\text{H}$  NMR

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and  $^{13}\text{C}$  NMR spectroscopy: m. p.  $78.5^\circ$ ;  $[\alpha]_D +31.7^\circ$  (c, 0.984,  $\text{CHCl}_3$ );  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  0.77 (3H), 0.91 (6H), 1.00-1.15 (2H), 1.40-2.10 (6H), 3.56 (1H), 3.82 (1H), 4.79 (1H), 5.62 (1H);  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ )  $\delta$  16.7, 21.2, 22.5, 23.8, 26.7, 30.0, 32.0, 34.6, 41.1, 47.6, 77.7, 168.1, 172.9.

Example 6

(1'R,2'S,5'R)-MENTHYL-5S-HYDROXY-1,3-OXATHIOLANE-2S-CARBOXYLATE, (1'R,2'S,5'R)-MENTHYL-5R-HYDROXY-1,3-OXATHIOLANE-2R-CARBOXYLATE, (1'R,2'S,5'R)-MENTHYL-5S-HYDROXY-1,3-OXATHIOLANE-2R-CARBOXYLATE, (1'R,2'S,5'R)-MENTHYL-5R-HYDROXY-1,3-OXATHIOLANE-2S-CARBOXYLATE



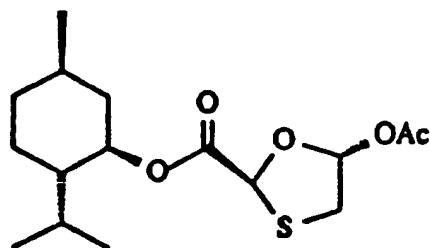
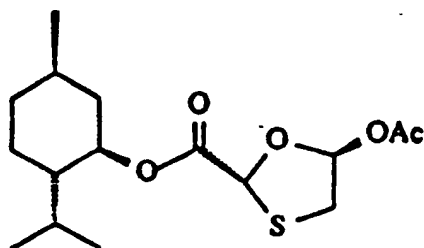
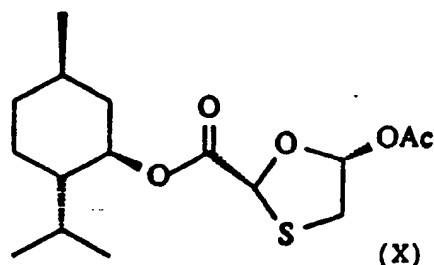
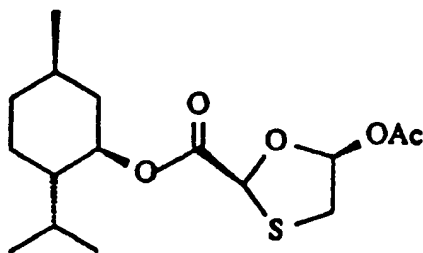
A freshly prepared solution of disiamylborane (13.4 mmol, 0.5 M in THF) was added via canula to a stirred solution of a 1:1 mixture of the menthyl ester carboxylate of formula (XVII) (1.28 g, 4.47 mmol) in THF (10 mL) at  $0^\circ\text{C}$  under an argon atmosphere. The resulting clear solution was stirred for 15 minutes at  $0^\circ\text{C}$  and 18 hours at ambient temperature. The reaction was quenched with methanol (5 mL), concentrated, and diluted with methylene chloride (20 mL). The resultant solution was washed with brine (5x2 mL) and dried over anhydrous magnesium sulfate. Removal of the solvent gave a clear oil. Subjecting this material to silica gel column chromatography (EtOAc-hexanes, 1:2, V/V) gave 0.65 g (50%) of the expected lactols in four diastereomeric forms:  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  0.71-2.09 (m,

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18H), 3.01-3.09 (m, 1H), 3.24-3.33 (m, 1H), 4.66-4.83 (m, 1H), 5.53-5.59 (m, 1H), 5.88-6.09 (m, 1H).

Example 7

5 (1'R,2'S,5'R)-MENTHYL-5S-ACETOXY-1,3-OXATHIOLANE-2S-CARBOXYLATE, (1'R,2'S,5'R)-MENTHYL-5R-ACETOXY-1,3-OXATHIOLANE-2R-CARBOXYLATE, (1'R,2'S,5'R)-MENTHYL-5S-ACETOXY-1,3-OXATHIOLANE-2R-CARBOXYLATE, (1'R,2'S,5'R)-MENTHYL-5R-ACETOXY-1,3-OXATHIOLANE-2S-CARBOXYLATE



10 The four title compounds were prepared as a mixture by the following two methods.

Method A

Lactols of formula (XVIII) (0.65 g, 2.25 mmol) were dissolved in anhydrous pyridine (1.5 mL) and  
 15 methylene chloride (5 mL). Acetyl chloride (0.5 mL, 7.0 mmol) was slowly added to this solution at 0°C. The resulting white suspension was stirred at ambient temperature for 3 hours. The reaction was then quenched with saturated aqueous ammonium chloride  
 20 solution (1 mL). The mixture was extracted with

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methylene chloride (5x2 mL) and the combined extract was concentrated to give a brown gummy material. This material was subjected to column chromatography (EtOAc-hexane, 1:3 V/V) to provide 0.3 g of the four acetates  
5 as a light yellow oil:  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  0.75 (d, 6H,  $J=7$  Hz), 0.78 (d, 6H,  $J=7$  Hz), 0.88-0.94 (m, 24H), 0.97-2.03 (m, 36H), 2.10 (s, 9H), 2.13 (s, 3H), 3.15 (d, 2H,  $J=12$  Hz), 3.23-3.30 (m, 4H), 3.42 (dd, 1H,  $J=4, 12$  Hz), 3.44 (dd, 1H,  $J=4, 12$  Hz), 4.65-4.75 (m, 4H), 5.61 (s, 1H),  
10 5.62 (s, 1H), 5.63 (s, 1H), 5.64 (s, 1H), 6.64 (m, 4H).

#### Method B

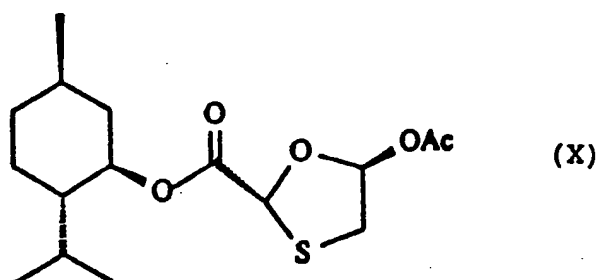
A solution of dicyclohexyl-carbodiimide (21.86 g, 0.106 mol) in dichloromethane (100 mL) was  
15 added to a 500 mL round bottom flask containing a solution of *trans*- and *cis*- 5-acetoxy-1,3-oxathiolane-2-carboxylic acid (X) (18.5 g, 0.096 mol), (1R,2S,5R)-(-)-menthol (16.5 g, 0.106 mol), and 4-dimethylaminopyridine (1.17 g, 9.63 mmol) in dichloromethane  
20 (200 mL) at 0°C. The resulting thick white slurry was stirred at room temperature for 3 hours at which time methanol (4.0 mL) and glacial acetic acid (2.0 mL) were added. After stirring for 10 minutes, the reaction mixture was diluted with hexanes (200 mL) and filtered  
25 through Celite. Subsequent removal of the solvent provided 32.5 g of the crude product. This substance was redissolved in hexanes (100 mL), filtered through Celite and concentrated to yield 30.5 g of material which was further purified by column chromatography  
30 (eluent: 100% hexanes to 5% EtOAc-hexanes) to give 5.5 g of a mixture (ca. 1:1) of (1'R,2'S,5'R)-menthyl-5R-acetoxy-1,3-oxathiolane-2S-carboxylate and (1'R,2'S,5'R)-menthyl-5S-acetoxy-1,3-oxathiolane-2R-carboxylate; 10.28 g of a material which contained

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mainly the above two diastereomers along with  
 (1'R,2'S,5'R)-menthyl-5S-acetoxy-1,3-oxathiolane-2S-  
 carboxylate and (1'R,2'S,5'R)-menthyl-5R-acetoxy-1,3-  
 oxathiolane-2R-carboxylate; 7.6 g of a random mixture  
 5 of the above four diastereomers; and 2.2 g of a mixture  
 (ca. 1:1) of (1'R,2'S,5'R)-menthyl-5S-acetoxy-1,3-  
 oxathiolane-2S-carboxylate and (1'R,2'S,5'R)-menthyl-  
 5R-acetoxy-1,3-oxathiolane-2R-carboxylate.

#### Example 8

10 (1'R,2'S,5'R)-MENTHYL-5R-ACETOXY-1,3-OXATHIOLANE-2R-  
CARBOXYLATE



(1'R,2'S,5'R)-Menthyl-5R-acetoxy-1,3-  
 oxathiolane-2R-carboxylate was prepared by the  
 15 following three methods.

#### Method A

A mixture of (1'R,2'S,5'R)-menthyl-5S-  
 acetoxy-1,3-oxathiolane-2S-carboxylate and  
 (1'R,2'S,5'R)-menthyl-5R-acetoxy-1,3-oxathiolane-2R-  
 20 carboxylate (5.5 g) obtained from Example 7 was  
 dissolved in petroleum ether (40-60°C) containing a  
 minimum amount of diethyl ether and cooled in a dry  
 ice-acetone bath. The white solid precipitate was  
 immediately collected by suction filtration to give 1.6  
 25 g of (1'R,2'S,5'R)-menthyl-5R-acetoxy-1,3-oxathiolane-  
 2R-carboxylate: m.p. 105.2°C;  $[\alpha]_D$  -60° (c, 0.51,

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CHCl<sub>3</sub>); <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 0.77 (d, 3H, J=7 Hz), 0.91 (d, 3H, J=7 Hz), 0.92 (d, 3H, J=7 Hz), 0.86-2.06 (m, 9H), 2.10 (s, 3H), 3.16 (d, 1H, J=12 Hz), 3.44 (dd, 1H, J=4, 12 Hz), 4.74 (dt, 1H, J=5, 12 Hz), 5.63 (s, 1H), 6.79  
5 (d, 1H, J=4 Hz); <sup>13</sup>C NMR (CDCl<sub>3</sub>) δ 16.16, 20.74, 21.11, 21.97, 23.29, 26.08, 31.38, 34.13, 37.24, 40.62, 47.07, 76.11, 79.97, 99.78, 168.60, 169.68.

#### Method B

A mixture of the four diastereomers of  
10 formula (X) (300 mg) was dissolved in n-pentane containing a minimum amount of diethyl ether and was kept at -20°C for 24 hours. The white needles formed were filtered quickly while cold to give 25 mg of material. The substance thus isolated was found to be  
15 identical in all respects with those obtained by Method A or C.

#### Method C

A solution of dicyclohexylcarbodiimide (1.362 g, 6.6 mmol) in dichloromethane (5 mL) was added to a  
20 50 mL round bottom flask containing a solution of trans-5-acetoxy-1,3-oxathiolane-2-carboxylic acid (1.16 g, 6.04 mmol), (1R,2S,5R)-(-)-menthol (1.038 g, 6.60 mmol), and 4-dimethylaminopyridine (75 mg, 0.62 mmol) in dichloromethane (10 mL) at 0°C. The resulting white  
25 slurry was stirred at room temperature for 3 hours at which time methanol (0.2 mL) and glacial acetic acid (0.2 mL) were added. After stirring for 10 minutes, the reaction mixture was diluted with hexanes (25 mL), filtered through Celite, and concentrated. The crude  
30 product thus obtained was dissolved in hexanes (25 mL), filtered through Celite and concentrated to provide 1.98 g (100%) of (1'R,2'S,5'R)-menthyl-5R-acetoxy-1,3-



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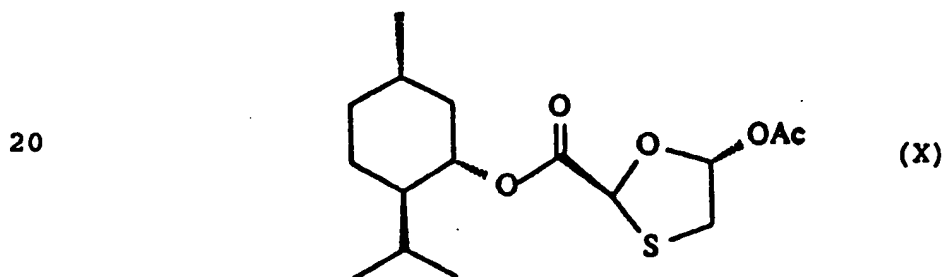
oxathiolane-2R-carboxylate and (1'R,2'S,5'R)-menthyl-5S-acetoxy-1,3-oxathiolane-2S-carboxylate:

<sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 0.75 (d, 3H, J=7 Hz), 0.78 (d, 3H, J=7 Hz), 0.85-0.92 (m, 12H), 0.95-2.19 (m, 18H), 2.10 (s, 6H), 3.15 (d, 2H, J=12 Hz), 3.42 (dd, 1H, J=4, 12 Hz), 3.44 (dd, 1H, J=4, 12 Hz), 4.74 (dt, 2H, J=5, 12 Hz), 5.61 (s, 1H), 5.62 (s, 1H), 6.65 (s, 2H)

The above mixture of diastereoisomers was dissolved in petroleum ether (40-60°C) containing a minimum amount of diethyl ether and was cooled in a dry ice-acetone bath. The white solid precipitate was immediately collected (620 mg) by suction filtration. This material was recrystallized again under the same conditions to yield 450 mg of a white solid. This compound was found to be identical in all respects to those prepared using either method A or method B.

#### Example 9

(1'S,2'R,5'S)- MENTHYL-5S-ACETOXY-1,3-OXATHIOLANE-2S-CARBOXYLATE



A solution of dicyclohexylcarbodiimide (491 mg, 2.38 mmol) in dichloromethane (7 mL) was added to a 50 mL round bottom flask containing a solution of trans-5-acetoxy-1,3-oxathiolane-2-carboxylic acid (IX) (416 mg, 2.2 mmol), (1S,2R,5S)-(+)-menthol (372 mg, 2.38 mmol), and 4-dimethylamino-pyridine (26 mg, 0.21

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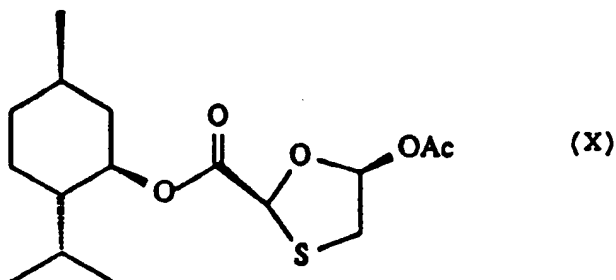
mmol) in dichloromethane (5 mL) at 0°C. The resulting thick slurry was stirred at room temperature for 3 hours at which time methanol (0.2 mL) and glacial acetic acid (0.2 mL) were added. After stirring for 10 minutes, the mixture was diluted with hexanes (25 mL), filtered through Celite, and concentrated. The crude product obtained was dissolved in hexanes (25 mL), filtered through Celite, and concentrated to produce 0.715 mg (100%) of two diastereomers, namely (1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolane-2S-carboxylate and (1'S,2'R,5'S)-menthyl-5R-acetoxy-1,3-oxathiolane-2R-carboxylate: <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 0.75 (d, 6H, J=7 Hz), 0.85-0.92 (m, 12H), 0.95-2.19 (m, 18H), 2.10 (s, 6H), 3.15 (d, 2H, J=12 Hz), 3.42 (dd, 1H, J=4, 12 Hz), 3.44 (dd, 1H, J=4, 12 Hz), 4.72 (dt, 2H, J=5, 12 Hz) 5.61 (s, 1H), 5.62 (s, 1H), 6.65 (s, 2H).

The above diastereomeric acetoxy menthyl esters mixture was dissolved in petroleum ether (40-60°C) containing a minimum amount of diethyl ether and was cooled in a dry ice-acetone bath. The white solid precipitate was immediately collected (200 mg) by suction filtration. This material was recrystallized again under the same conditions to yield 130 mg (34% based on one enantiomer) of (1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolane-2S-carboxylate: m.p. 104.2°C; [α]<sub>D</sub> +59.2° (c, 1.02, CHCl<sub>3</sub>); <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 0.77 (d, 3H, J=7 Hz), 0.91 (d, 3H, J=7 Hz), 0.92 (d, 3H, J=7 Hz), 0.86-2.06 (m, 9H), 2.10 (s, 3H), 3.16 (d, 1H, J=12 Hz), 3.44 (dd, 1H, J=4, 12 Hz), 4.74 (dt, 1H, J=5, 12 Hz), 5.63 (s, 1H), 6.79 (d, 1H, J=4 Hz); <sup>13</sup>C NMR (CDCl<sub>3</sub>) δ 16.16, 20.74, 21.11, 21.97, 23.29, 26.08, 31.38, 34.13, 37.24, 40.62, 47.07, 76.11, 79.96, 99.78, 168.60, 169.68.

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Example 10

(1'R,2'S,5'R)-MENTHYL-5R-ACETOXY-1,3-OXATHIOLANE-2S-CARBOXYLATE



5 (1'R,2'S,5'R)-Methyl-5R-acetoxy-1,3-oxathiolane-2S-carboxylate was prepared by the following two methods.

Method A

A saturated solution of a mixture of the four  
 10 diastereomers (12.28 g), obtained in Example 7, was prepared in petroleum ether containing a minimum amount of diethyl ether and was kept at -20°C for 72 hours. The white crystalline solid produced was isolated by filtration to give 1.6 g of (1'R,2'S,5'R)-menthyl-5R-  
 15 acetoxy-1,3-oxathiolane-2S-carboxylate: m.p. 110.2°C;  $[\alpha]_D -177^\circ$  (c, 0.7, CHCl<sub>3</sub>); <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 0.75 (d, 3H, J=7 Hz), 0.88 (d, 3H, J=7 Hz), 0.92 (d, 3H, J=7 Hz), 0.97-2.02 (m, 9H), 2.12 (s, 3H), 3.22 (d, 1H, J=11 Hz), 3.29 (dd, 1H J=4, 11 Hz), 4.74 (dt, 1H, J=4, 11  
 20 Hz), 5.63 (s, 1H), 6.65 (d, 1H, J=3 Hz); <sup>13</sup>C NMR (CDCl<sub>3</sub>) δ 16.9, 20.69, 21.19, 21.95, 23.29, 26.10, 31.34, 34.0, 37.62, 40.32, 46.82, 75.69, 80.20, 99.36, 168.55, 170.23.

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Method B

A solution of dicyclohexylcarbodiimide (118 mg, 0.572 mmol) in dichloromethane (5 mL) was added to a 25 mL round bottom flask containing a solution of

5 cis-5-acetoxy-1,3-oxathiolane-2-carboxylic acid (100 mg, 0.52 mmol), (1R,2S,5R)-(-)-menthol (85 mg, 0.54 mmol), and 4-dimethyl-aminopyridine (DMAP) (8 mg, 0.053 mmol) in dichloromethane (10 mL) at 0°C. The resulting white slurry was stirred at room temperature for 3

10 hours at which time methanol (0.1 mL) and glacial acetic acid (0.1 mL) was added. After stirring for 10 minutes, the mixture was diluted with hexanes (15 mL), filtered through Celite, and concentrated. The crude product obtained was dissolved in hexanes (15 mL),

15 filtered through Celite, and concentrated to yield 170 mg (100%) of (1'R,2'S,5'R)-menthyl-5R-acetoxy-1,3-oxathiolane-2S-carboxylate and (1'R,2'S,5'R)-menthyl-5S-acetoxy-1,3-oxathiolane-2R-carboxylate: <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 0.75 (d, 3H, J=7 Hz), 0.78 (d, 3H, J=7 Hz),

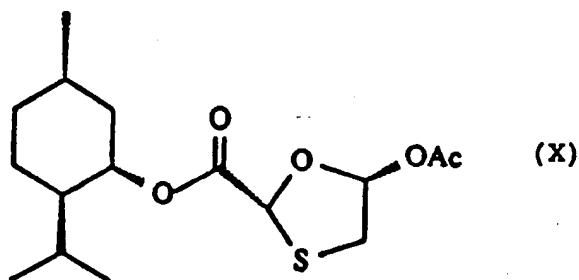
20 0.88-0.94 (m, 12H), 0.97-2.03 (m, 18H), 2.10 (s, 3H), 2.13 (s, 3H), 3.23-3.30 (m, 4H), 4.65-4.75 (m, 2H), 5.63 (s, 1H), 5.64 (s, 1H), 6.64 (m, 2H).

The above mixture of diastereomers was recrystallized from petroleum ether (40-60°C) and a

25 minimum amount of diethyl ether at room temperature. The white crystalline material formed was collected (95 mg) by filtration. This material was recrystallized again from diethyl ether-petroleum ether to yield 74 mg (78% based on one enantiomer) of

30 (1'R,2'S,5'R) menthyl-5R-acetoxy-1,3-oxathiolane-2S-carboxylate.

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Example 11(1'S,2'R,5'S)-MENTHYL-5S-ACETOXY-1,3-OXATHIOLANE-2R-CARBOXYLATE

5                   A solution of dicyclohexylcarbodiimide (1.588 g, 7.7 mmol) in dichloromethane (7 mL) was added to a 50 ml round bottom flask containing a solution *cis*-5-acetoxy-1,3-oxathiolane-2-carboxylic acid (1.36 g, 7 mmol), (1S,2R,5S)-(+)-menthol (1.216 g, 7.7 mmol), and  
10 4-dimethylamino-pyridine (85 mg, 0.7 mmol) in dichloromethane (16 mL) at 0°C. The resulting thick slurry was stirred at room temperature for 3 hours. The reaction was quenched with methanol (0.4 mL) and glacial acetic acid (0.4 mL) and the mixture was  
15 stirred for 10 min. The resultant mixture was diluted with hexanes (25 mL), filtered through a pad of Celite, and concentrated. The crude material thus obtained was redissolved in hexanes (25 mL) and filtered through Celite. Removal of the solvent under reduced pressure  
20 yielded 2.3 g of a white solid (100%) which consisted of (1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolane-2R-carboxylate and (1'S,2'R,5'S)-menthyl-5R-acetoxy-1,3-oxathiolane-2S-carboxylate:

<sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 0.75 (d, 3H, J=7 Hz), 0.78 (d, 3H, J=7 Hz), 0.88-0.94 (m, 12H), 0.97-2.03 (m, 18H), 2.10 (s, 3H), 2.13 (s, 3H), 3.23-3.30 (m, 4H), 4.65-4.74 (m, 2H), 5.63 (s, 1H), 5.64 (s, 1H), 6.64 (m, 2H).

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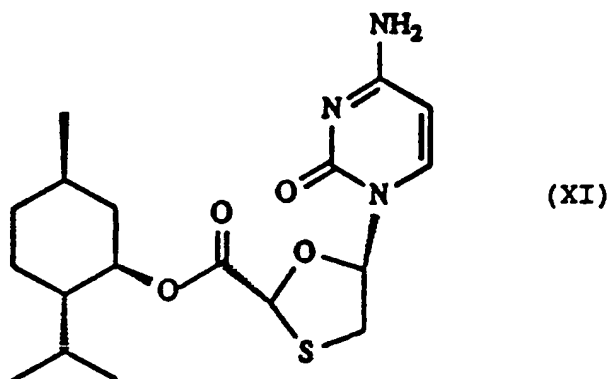
The above mixture of diastereomers was recrystallized from petroleum ether (40-60°C) and a small amount of diethyl ether at room temperature to give 1.3 g of a white solid. This material was

5 recrystallized again from diethyl ether-petroleum ether (40-60°C) to give 900 mg (78% based on one enantiomer) of (1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolane-2R-carboxylate: m.p. 110.2°C;  $[\alpha]_D +177^\circ$  (c, 1.0, CHCl<sub>3</sub>); <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 0.75 (d, 3H, J=7 Hz), 0.89 (d, 3H, J=7 Hz), 0.92 (d, 3H, J=7 Hz), 0.98-2.02 (m, 9H), 2.12 (s, 3H), 3.22 (d, 1H, J=11 Hz), 3.29 (dd, 1H, J=4, 11 Hz), 4.74 (dt, 1H, J=11, 4 Hz), 5.63 (s, 1H), 6.65 (d, 1H, J=3 Hz); <sup>13</sup>C NMR (CDCl<sub>3</sub>) δ 16.9, 20.69, 21.19, 21.95, 23.29, 26.10, 31.34, 34.09, 37.62, 40.32, 46.82,

15 75.79, 80.20, 99.36, 168.55, 170.23.

Example 12

(1'R,2'S,5'R)-MENTHYL-5S-(CYTOSIN-1"-YL)-1,3-OXATHIOLANE-2R-CARBOXYLATE



20 t-Butyl-dimethylsilyl trifluoromethane-sulfonate (1.1 mL, 4.79 mmol) was added to a suspension of cytosine (0.27 g, 2.5 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (2 mL) containing 2,4,6-collidine (0.65 ml, 4.92 mmol) at room temperature. The resultant mixture was stirred for 15

25 minutes and a clear solution was produced. A solution

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of (1'R,2'S,5'R)-menthyl-5S-acetoxy-1,3-oxathiolane-2R-carboxylate (0.66 g, 1.99 mmol) in methylene chloride (1.5 mL) was added to the mixture and stirring was continued for 5 minutes. Iodotrimethylsilane

5 (0.31 mL, 2.18 mmol) was introduced dropwise and a white precipitate was produced when the addition was completed. The reaction mixture was allowed to stir for 18 hours. The reaction was quenched by addition of a saturated aqueous solution of  $\text{Na}_2\text{S}_2\text{O}_3$  (10 mL) and

10  $\text{CH}_2\text{Cl}_2$  (30 mL). The organic layer was separated and washed with brine (2x10 mL). The solvent was removed in vacuo to give a viscous oil which was suspended in diethyl ether (30 mL). To this suspension was added a saturated aqueous solution of  $\text{NaHCO}_3$  (20 mL) with

15 vigorous stirring. A white precipitate appeared and the resultant suspension was diluted with hexanes (10 mL). The precipitate was collected by filtration to give 0.57 g (75%) of a white solid. The  $^1\text{H}$  NMR spectrum of this material indicated that it was a

20 mixture of the *cis*- and *trans*- diastereomers of the expected nucleoside in a 23:1 ratio.

This product was purified further by recrystallization from EtOAc-hexanes-MeOH:  $[\alpha]_D -144^\circ$  (c, 1.02,  $\text{CHCl}_3$ ); m.p.  $219^\circ\text{C}$  (decomposed);  $^1\text{H}$  NMR

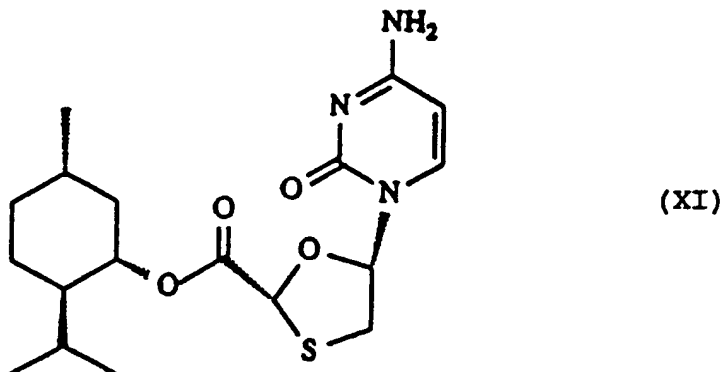
25 ( $\text{CDCl}_3$ )  $\delta$  0.76 (d, 3H,  $J=7$  Hz), 0.85-0.94 (m, 6H), 1.02-1.10 (m, 2H), 1.42-2.06 (m, 7H), 3.14 (dd, 1H,  $J=6.6, 12.1$  Hz), 3.54 (dd, 1H,  $J=4.7, 12.1$  Hz), 4.72-4.78 (m, 1H), 5.46 (s, 1H), 5.99 (d, 1H,  $J=7.5$  Hz), 8.43 (d, 1H,  $J=7.6$  Hz);  $^{13}\text{C}$  ( $\text{CDCl}_3$ )  $\delta$  16.1, 20.7, 21.9,

30 23.2, 26.4, 31.4, 34.0, 36.3, 40.7, 47.1, 76.7, 78.4, 90.3, 94.6, 141.8, 155.4, 165.6, 169.8.

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Example 13

(1'S,2'R,5'S)-MENTHYL-5S-(CYTOSIN-1"-YL)-1,3-OXATHIOLANE-2R-CARBOXYLATE



5                    2,4,6-Collidine (0.317 mL, 2.4 mmol) and t-butyltrimethylsilyl trifluoromethanesulfonate (0.551 mL, 2.4 mmol) were added successively to a suspension of cytosine (133.3 mg, 1.2 mmol) in  $\text{CH}_2\text{Cl}_2$  (1 mL) at room temperature under an argon atmosphere. The resultant

10 mixture was stirred for 15 minutes to produce a clear solution. A solution of (1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolane-2R-carboxylate (330 mg, 1 mmol) in  $\text{CH}_2\text{Cl}_2$  (0.5 mL) was introduced, followed by iodotrimethylsilane (0.156 mL, 1.1 mmol). The

15 resultant mixture was stirred for 3 hours. The mixture was diluted with  $\text{CH}_2\text{Cl}_2$  (20 mL) and washed successively with saturated aqueous  $\text{NaHSO}_3$ , water, and brine. The solvent was evaporated and the residue was taken up in ether-hexanes (1:1, 10 mL) and saturated aqueous  $\text{NaHCO}_3$

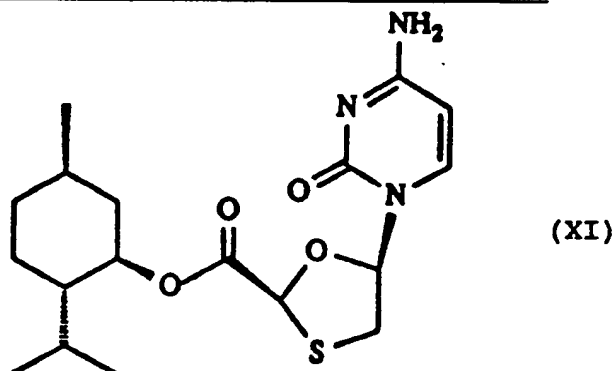
20 (2 mL). Stirring was continued for 15 minutes. The aqueous layer was removed and the organic phase was centrifuged to give a white solid which was washed with hexanes (3x5 mL) and dried under vacuum. This substance, namely (1'S,2'R,5'S)-menthyl-5S-(cytosin-

25 1"-yl)-1,3-oxathiolan-2R-carboxylate (380 mg, 100%) was contaminated with about 3% of (1'S,2'R,5'S)-menthyl-



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5R-(cytosin-1"-1,3-oxathiolan-2R-carboxylate (as indicated by its  $^1\text{H}$  NMR spectrum), was recrystallized from MeOH to give (1'S,2'R,5'S)-menthyl-5S-(cytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate:  $[\alpha]_D -58^\circ$  (c, 0.506,  $\text{CHCl}_3$ ); m.p.:  $235^\circ\text{C}$  (decomposed));  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  0.80 (3H), 0.92 (6H), 1.06 (2H), 1.37-2.10 (7H), 3.11 (1H), 3.55 (1H), 4.77 (1H), 5.47 (1H), 5.79 (1H), 6.49 (1H), 8.37 (1H);  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ )  $\delta$  6.8, 21.3, 22.5, 23.9, 26.8, 32.0, 34.6, 37.0, 40.7, 47.4, 77.3, 79.3, 90.9, 95.3, 142.9, 155.1, 164.9, 170.1.

Example 14(1'R,2'S,5'R)-MENTHYL-5R-(CYTOSIN-1"-YL)-1,3-OXATHIOLANE-2S-CARBOXYLATE

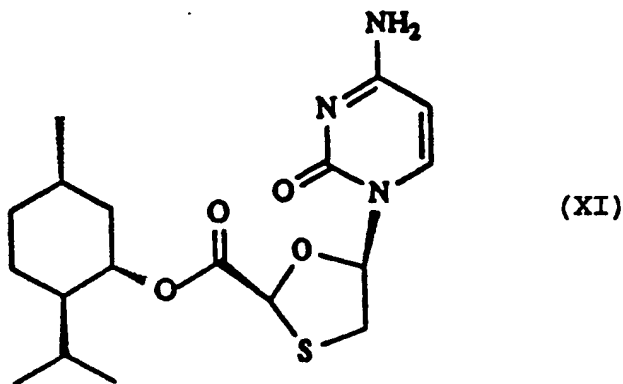
15                    2,4,6-collidine (0.317 mL, 2.4 mmol) and t-butyldimethylsilyl trifluoromethanesulfonate (0.551 mL, 2.4 mmol) were added successively to a suspension of cytosine (133.3 mg, 1.2 mmol) in  $\text{CH}_2\text{Cl}_2$  (1 mL) at room temperature under an argon atmosphere. The resultant  
20 mixture was stirred for 15 minutes and a clear solution was obtained. A solution of (1'R,2'S,5'R)-menthyl-5R-acetoxy-1,3-oxathiolan-2S-carboxylate (330 mg, 1 mmol) in  $\text{CH}_2\text{Cl}_2$  (0.5 mL) was introduced, followed by iodotrimethylsilane (0.156 mL, 1.1 mmol). Stirring was  
25 continued for 3 hours. The mixture was diluted with  $\text{CH}_2\text{Cl}_2$  (20 mL) and washed successively with saturated aqueous  $\text{NaHSO}_3$ , water, brine and then was concentrated.

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The residue was taken up in ether-hexanes (1:1, 10 mL) and saturated aqueous  $\text{NaHCO}_3$  (2 mL) and was stirred at room temperature for 15 minutes. The aqueous layer was removed and the organic phase was centrifuged to yield  
5 a white solid which was washed with hexanes (3x5 mL) and then dried under vacuum. The product (1'R,2'S,5'R)-menthyl-5R-(cytosin-1"-yl)-1,3-oxathiolan-2S-carboxylate (336.3 mg, 88%) contained about 6% of (1'R,2'S,5'R)-menthyl-5S-(cytosin-1"-yl)-  
10 1,3-oxathiolan-2S-carboxylate (NMR). This material was recrystallized from MeOH to give the desired product:  $[\alpha]_D^{+56}$  (c, 1.08,  $\text{CHCl}_3$ ); m.p.: 235°C (decomposed);  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  0.80 (3H), 0.91 (6H), 1.00 (2H), 1.37-2.10 (7H), 3.11 (1H), 3.55 (1H), 4.77 (1H), 5.47 (1H),  
15 5.79 (1H), 6.49 (1H), 8.37 (1H);  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ )  $\delta$  16.8, 21.3, 22.5, 23.9, 26.8, 32.0, 34.6, 36.8, 40.7, 47.4, 77.1, 78.8, 90.9, 95.6, 141.9, 156.3, 166.6, 170.2.

#### Example 15

20 (1'S,2'R,5'S)-MENTHYL-5R-(CYTOSIN-1"-YL)-1,3-OXATHIOLANE-2S-CARBOXYLATE



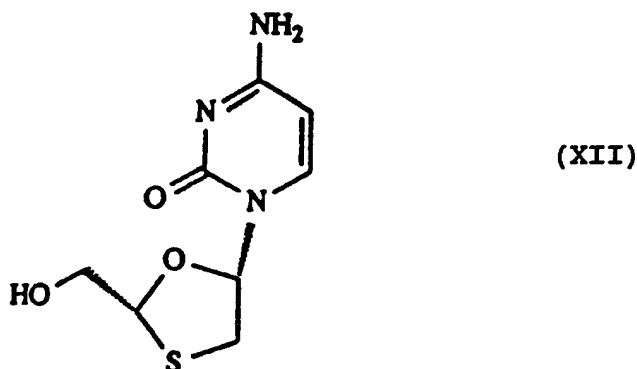
2,4,6-collidine (0.106 mL, 0.8 mmol) and t-butyltrimethylsilyl trifluoromethanesulfonate were added successively to a suspension of cytosine (44 mg, 0.4  
25 mmol) in  $\text{CH}_2\text{Cl}_2$  (0.5 mL) at room temperature under an argon atmosphere. Stirring was continued at room

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temperature for 15 minutes and a clear solution was produced. A solution of (1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolan-2S-carboxylate (110 mg, 0.33 mmol) in  $\text{CH}_2\text{Cl}_2$  (0.3 mL) was added, followed by  
5 iodotrimethylsilane (0.052 mL, 0.36 mmol). The resultant mixture was stirred at room temperature overnight and then was diluted with  $\text{CH}_2\text{Cl}_2$  (10 mL). The mixture was washed successively with saturated aqueous  $\text{NaHSO}_3$ , water, brine and concentrated under reduced  
10 pressure. The residue was taken up in ether-hexanes (1:1, 5 mL) and saturated aqueous  $\text{NaHCO}_3$  (1 mL) and stirring was continued at room temperature for 20 minutes. The aqueous layer was removed and the white solid suspended in the organic phase was collected by  
15 centrifugation. This solid was washed with hexanes (3x5 mL) and dried under vacuum to provide 65 mg (51.2%) of (1'S,2'R,5'S)-menthyl-5R-(cytosin-1"-yl)-1,3-oxathiolan-2S-carboxylate contaminated with approximately 5% of (1'S,2'R,5'S)-menthyl-5S-(cytosin-  
20 1"-yl)-1,3-oxathiolan-2S-carboxylate as indicated by  $^1\text{H}$  NMR spectroscopy. Recrystallization of the crude material from  $\text{MeOH-Et}_2\text{O}$  gave the desired product:  
m.p. 210-211°C;  $[\alpha]_D +179^\circ$  (c, 0.66,  $\text{CHCl}_3$ );  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  0.77 (3H) 0.92 (6H), 1.00 (2H), 1.37-2.10  
25 (6H), 3.14 (1H), 3.55 (1H), 4.76 (1H), 5.46 (1H), 5.88 (1H), 6.46 (1H), 8.38 (1H);  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ )  $\delta$  16.8, 21.3, 21.8, 22.5, 23.9, 26.7, 31.9, 34.7, 38.7, 40.9, 47.4, 76.4, 80.8, 100.0, 169.1, 170.8

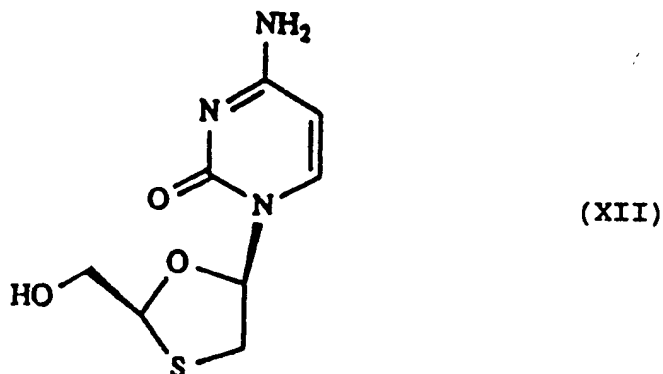
The washings and the supernatant were  
30 combined and washed with 1N  $\text{HCl}$ , water, brine, and then was dried over  $\text{Na}_2\text{SO}_4$ . Evaporation of the solvent yielded 53 mg (48%) of unreacted (1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolan-2S-carboxylate.

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Example 162R-HYDROXYMETHYL-5S-(CYTOSIN-1'-YL)-1,3-OXATHIOLANE

A solution of (1'R,2'S,5'R)-menthyl-5S-(cytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate (67 mg, 0.18 mmol) in THF (1 mL) was slowly added to a stirred suspension of lithium aluminum hydride (19 mg, 0.5 mmol) in THF (2 mL) at ambient temperature under an argon atmosphere. Stirring was continued for 30 minutes. The reaction was then quenched with methanol (3 mL), followed by the addition of silica gel (5 g). The resultant slurry was stirred for 30 minutes and then was transferred to a short column packed with Celite and silica gel and was eluted with a 1:1:1 mixture of EtOAc-hexane-methanol (50 mL). The eluate was concentrated and subjected to silica gel column chromatography (EtOAc-hexane-methanol, 1:1:1) to give a gummy solid. This solid was dried azeotropically with toluene to give 38 mg (94%) of the desired product:

[ $\alpha$ ]<sub>D</sub> -122° (c, 1.01, MeOH); m.p. 128-130°C; <sup>1</sup>H NMR (CD<sub>3</sub>OD)  $\delta$  3.05 (dd, 1H, J=4.3, 11.9 Hz) 3.42 (dd, 1H, J=5.3, 11.9 Hz), 3.76-3.89 (m, 2H), 5.19-5.21 (m, 1H), 5.81 (d, 1H, J=7.6 Hz), 6.20-6.23 (m, 1H), 7.01-7.16 (brm, 2H, exchangeable), 7.98 (d, 1H, J=7.5 Hz); <sup>13</sup>C (CD<sub>3</sub>OD)  $\delta$  38.5, 64.1, 88.0, 88.9, 95.7, 142.8, 157.9, 167.7.

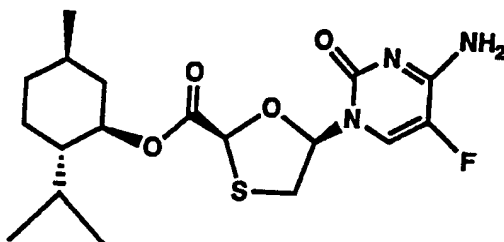
Example 172S-HYDROXYMETHYL-5R-(CYTOSIN-1'-YL)-1,3-OXATHIOLANE

A solution of (1'R,2'S,5'R)-menthyl-5R-  
 5 (cytosin-1"-yl)-1,3-oxathiolane-2S-carboxylate (102 mg,  
 0.27 mmol) in THF (3 mL) was slowly added to a stirred  
 suspension of lithium aluminum hydride (20 mg, 0.54  
 mmol) in THF (2 mL) at ambient temperature under an  
 argon atmosphere. Stirring was continued for 30  
 10 minutes and the reaction was quenched with methanol  
 (5 mL), followed by the addition of silica gel (7 g).  
 The resultant slurry was stirred for 30 minutes,  
 transferred to a short column packed with Celite and  
 silica gel and was eluted with a 1:1:1 mixture of  
 15 EtOAc-hexane-MeOH (50 mL). The eluate was concentrated  
 and subjected to silica gel column chromatography  
 (EtOAc-hexane-MeOH, 1:1:1) to provided a gummy solid  
 which was dried azeotropically with toluene to give  
 50 mg (82%) of a white solid as the product:  $[\alpha]_D$   
 20  $+125^\circ$  (c, 1.01, MeOH); m.p. 130-132°C;  $^1\text{H}$  NMR ( $\text{CD}_3\text{OD}$ )  $\delta$   
 3.05 (dd, 1H,  $J=4.3, 11.9$  Hz), 3.42 (dd, 1H,  $J=5.3,$   
 11.9 Hz), 3.76-3.89 (m, 2H), 5.19-5.21 (m, 1H), 5.81  
 (d, 1H,  $J=7.6$  Hz), 6.20-6.23 (m, 1H), 7.01-7.16 (brm,  
 2H, exchangeable), 7.98 (d, 1H,  $J=7.5$  Hz);  $^{13}\text{C}$  ( $\text{CD}_3\text{OD}$ )  $\delta$   
 25 38.5, 64.1, 88.0, 88.9, 95.7, 142.8, 157.9, 167.7.

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Example 18

(1'R,2'S,5'R)-MENTHYL-5R-(5'-FLUOROCYTOSIN-1"-YL)-1,3-OXATHIOLANE-2S-CARBOXYLATE



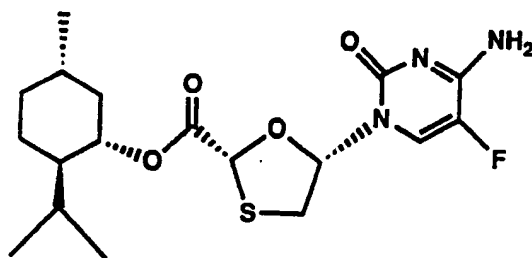
To a suspension of 5-fluorocytosine (155 mg,  
5 1.2 mmol) in  $\text{CH}_2\text{Cl}_2$  (1 mL) at room temperature under an  
argon atmosphere was added, successively, 2,4,6-  
collidine (0.317 mL, 2.4 mmol) and t-butyldimethylsilyl  
trifluoromethane-sulfonate (0.551 mL, 2.4 mmol). The  
resultant mixture was stirred for 15 minutes and a  
10 clear solution was obtained. A solution of  
(1'R,2'S,5'R)-menthyl-5R-acetoxy-1,3-oxathiolane-2S-  
carboxylate (330 mg, 1 mmol) in  $\text{CH}_2\text{Cl}_2$  (0.5 mL) was  
introduced, followed by iodotrimethylsilane (0.156 mL,  
1.1 mmol). Stirring was continued for 3 hours. The  
15 mixture was diluted with  $\text{CH}_2\text{Cl}_2$  (20 mL) and washed  
successively with saturated aqueous  $\text{NaHSO}_3$ , water,  
brine and then was concentrated. The residue was taken  
up in ether-hexanes (1:1, 10mL) and saturated aqueous  
 $\text{NaHCO}_3$  (2 mL) and stirred at room temperature for 15  
20 minutes. The aqueous layer was removed and the organic  
phase was centrifuged to afford a white solid which was  
washed with hexanes (3x5 mL) and then dried under  
vacuum. The product (1'R,2'S,5'R)-menthyl-5R-(5"-  
fluorocytosin-1"-yl)-1,3-oxathiolane-2S-carboxylate  
25 (350 mg, 88%) thus obtained contained about 6% of  
(1'R,2'S,5'R)-menthyl-5S-(5"-fluorocytosin-1"-yl)-1,3-

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oxathiolane-2S-carboxylate (NMR). This material was recrystallized from MeOH/CH<sub>2</sub>Cl<sub>2</sub>/benzene to give a crystalline product:  $[\alpha]_D^{26} +22^\circ$  (c, 0.19, MeOH); m.p. 216-218°C, <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 0.78 (d, 3H, J= 7Hz), 0.91  
5 (t, 6H, J=7.3 Hz), 1.00 (m, 2H), 1.39-2.04 (m, 7H), 3.12 (dd, 1H, J=6.6 Hz, 6.1 Hz), 3.52 (dd, 1H, J=4.7 Hz, 6.1 Hz), 4.79 (dt, 1H, J=4.4 Hz, 4.3 Hz), 5.46 (s, 1 H), 5.75 (bs, 1H, exchangeable), 6.42 (5t, 1H, J=5.0 Hz), 8.10 (bs, 1H, exchangeable), 8.48 (d, 1H, J=6.6  
10 Hz); <sup>13</sup>C NMR (CDCl<sub>3</sub>-DMSO-d<sub>6</sub>): δ 16.7, 21.2, 22.4, 23.7, 26.6, 31.8, 34.4, 36.6, 40.5, 47.2, 77.1, 79.1, 90.8, 126.3 (d, J=33 Hz), 137.1 (d, J=244 Hz), 154.2, 158.3 (d, J=15 Hz), 170.1.

#### Example 19

15 (1'S,2'R,5'S)-MENTHYL-5S-(5"-FLUOROCYTOSIN-1"-YL)-1,3-OXATHIOLANE-2R-CARBOXYLATE



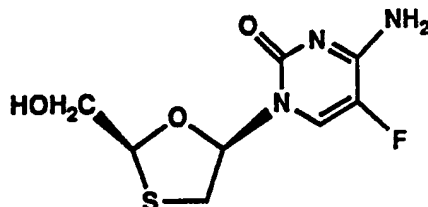
To a suspension of 5-fluorocytosine (180.0 mg, 1.4 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (1 mL) at room temperature under an argon atmosphere was added, successively, 2,4,6-  
20 collidine (0.46 mL, 3.5 mmol) and t-butyldimethylsilyl trifluoromethane-sulfonate (0.67 mL, 2.9 mmol). The resultant mixture was stirred for 15 minutes and a clear solution was obtained. A solution of (1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolane-2R-

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carboxylate (414 mg, 1.25 mmol) in  $\text{CH}_2\text{Cl}_2$  (0.6 mL) was introduced, followed by iodotrimethylsilane (0.18 mL, 1.27 mmol). The resultant mixture was stirred for 1 hour. The mixture was diluted with  $\text{CH}_2\text{Cl}_2$  (20 mL) and  
5 washed successively with saturated aqueous  $\text{NaHSO}_3$ , water, and brine. The solvent was evaporated and the residue was taken up in ether-hexanes (1:1, 10 mL) and saturated aqueous  $\text{NaHCO}_3$  (2 mL). Stirring was continued for 15 minutes. The aqueous layer was  
10 removed and the organic phase was centrifuged to give a white solid which was washed with hexanes (3x5 mL) and dried under vacuum. This substance, namely (1'S,2'R,5'S)-menthyl-5S-(5"-fluorocytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate (454 mg, 91%) contained  
15 about 7% of (1'S,2'R,5'S)-menthyl-5R-(5"-fluorocytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate (as indicated by its  $^1\text{H}$  NMR spectrum), was recrystallized from benzene  $\text{CH}_2\text{Cl}_2$ -MeOH to give the title compound:  $[\alpha]_D^{26} -20^\circ$  (c, 0.072, MeOH); m.p. 220-222°C (decomposed),  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  
20  $\delta$  0.80 (d, 3H,  $J=7$  Hz), 0.90 (t, 6H,  $J=7$  Hz), 1.0 (m, 2H), 1.39-2.04 (m, 7H), 3.12 (dd, 1H,  $J=6.6$  and 6 Hz), 3.52 (dd, 1H,  $J=5$  and 6 Hz), 4.8 (dt, 1H,  $J=4.4$  and 4.3 Hz), 5.46 (s, 1H), 5.78 (bs, 1H, exchangeable), 6.42 (t, 1H,  $J=5$  Hz), 8.1 (bs, 1H exchangeable), 8.5 d, 1H,  
25  $J=6.6$  Hz);  $^{13}\text{C}$  ( $\text{CDCl}_3$ )  $\delta$  16.2, 20.7, 21.9, 23.3, 26.2, 31.4, 34.0, 36.3, 40.1, 46.8, 76.7, 78.7, 90.5, 125.9 (d,  $J=33$  Hz), 136.5 (d,  $J=242$  Hz), 153.7, 158.2 (d,  $J=14$  Hz), 169.6.

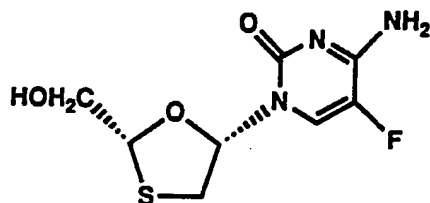


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Example 202S-HYDROXYMETHYL-5R-(5'-FLUOROCYTOSIN-1'-YL)-1,3-OXATHIOLANE

To a suspension of lithium aluminum hydride  
5 (10 mg, 0.54 mmol) in THF (1 mL) at ambient temperature  
under an argon atmosphere was slowly added a solution  
of (1'R,2'S,5'R)-menthyl-5R-(5"-fluorocytosin-1"-yl)-  
1,3-oxathiolane-2S-carboxylate (54 mg, 0.135 mmol) in  
THF (2 mL). The reaction mixture was allowed to stir  
10 for 30 minutes, then quenched with excess methanol (2  
mL), followed by the addition of silica gel (3 g). The  
resultant slurry was subjected to silica gel column  
chromatography (EtOAc-Hexane-MeOH, 1:1:1) to provide a  
gummy solid which was dried azeotropically with toluene  
15 to give 20.7 mg (63%) of a white solid as the product:  
[ $\alpha$ ]<sub>D</sub><sup>26</sup>+114° (c, 0.12, MeOH); <sup>1</sup>H NMR (DMSO-d<sub>6</sub>)  $\delta$  3.14  
(dd, 1H, J=4.3, 11.9 Hz), 3.42 (dd, 1H J=5.3, 11.9 Hz),  
3.76 (m, 2H), 5.18 (m, 1H), 5.42 (t, 1H, J=4.8 Hz), 6.14  
(m, 1H), 7.59 (br m, 1H, exchangeable), 7.83 (br m, 1H  
20 exchangeable), 8.20 (d, 1H, J=7.66 Hz).

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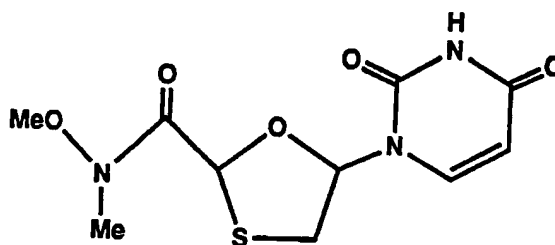
Example 212R-HYDROXYMETHYL-5S-(5'-FLUOROCYTOSIN-1'-YL)-1,3-  
OXATHIOLANE

To a stirred THF (2 mL) suspension of lithium  
5 aluminum hydride (22 mg, 1.13 mmol) at ambient  
temperature under an argon atmosphere was slowly added  
a solution of (1'R, 2'S, 5'R)-menthyl-5S-(5"-  
fluorocytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate (91  
mg, 0.23 mmol) in THF (8 mL). The reaction mixture was  
10 allowed to stir for 2 hours., and was quenched by  
addition of methanol (3 mL), followed by silica gel (5  
g). The resultant slurry was stirred for 30 minutes.  
The mixture was then passed through a short pad of  
Celite and silica gel eluted with a 1:1:1 mixture of  
15 EtOAc-hexane-Methanol (10x5 mL). The eluate was  
concentrated and subjected to silica gel column  
chromatography (EtOAc-hexane-methanol, 1:1:1) to give a  
gummy solid. This solid was dried azeotropically with  
toluene to give 45 mg (80%) of the desired product:  
20  $[\alpha]_D^{26} -119^\circ$  (c, 1.01, MeOH),  $^1\text{H}$  NMR (DMSO-d<sub>6</sub>)  $\delta$  3.14  
(dd, 1H, J=4.3, 11.9 Hz), 3.42 (dd, 1H, J=5.3, 11.9  
Hz), 3.76 (m, 2H), 5.18 (m, 1H), 5.42 (t, 1H, J=4.8  
Hz), 6.14 (m, 1H), 7.59 (br m, 1H, exchangeable), 7.83  
(br m, 1H exchangeable), 8.20 (d, 1H J=7.66 Hz).

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Example 22

CIS-2 (N-METHYL-N-METHOXYAMINOCARBONYL) -5-(URACIL-1'-YL) -1,3-OXATHIOLANE



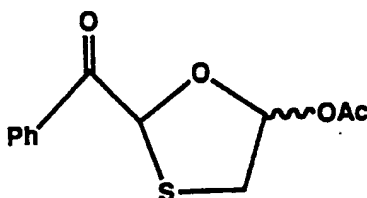
Trimethylsilyl trifluoromethanesulphonate  
5 (TMSOTf) (107  $\mu$ L, 0.552 mmol) was introduced to a stirred suspension of uracil (31 mg, 0.276 mmol) in dichloromethane (1.5 mL) containing collidine (73  $\mu$ L, 0.552 mmol) under argon atmosphere. The resultant mixture was stirred for 15 minutes to provide a  
10 homogeneous solution. A solution of *trans*-2-(N-methyl-N-methoxyaminocarbonyl)-5-acetoxy-1,3-oxathiolane (50 mg, 0.23 mmol) in dichloromethane (1mL) was introduced, followed by iodotrimethylsilane (TMSI) (33  $\mu$ L, 0.23 mmol). The reaction was allowed to proceed for 2.5  
15 hours and then was quenched with a solution of saturated  $\text{NaHCO}_3$  and  $\text{Na}_2\text{S}_2\text{O}_3$  (1:1). The resulting mixture was stirred for 5 minutes and then was transferred to a separatory funnel with the aid of more dichloromethane. The aqueous phase was removed and the  
20 organic layer was washed with saturated  $\text{Na}_2\text{S}_2\text{O}_3$ , water, brine and then was dried ( $\text{Na}_2\text{SO}_4$ ). Evaporation of the solvent under reduced pressure afforded the crude product which was triturated with EtOAc-Hexane (1:1) to give 54 mg (87%) of the title compound as a white  
25 solid;  $^1\text{H}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  3.14 (d of d, 1H,  $J=8.0$ , 11.8 Hz), 3.23 (s, 3H), 3.38 (d of d, 1H,  $J=4.7$ , 11.8 Hz),

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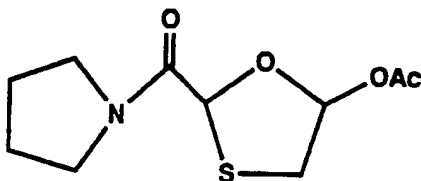
3.74 (s, 3H), 5.80 (d, 1H, J=8.2 Hz), 5.82 (s, 1H),  
6.44 (d of d, 1H, J=4.7, 8.0 Hz), 8.64 (d, 1H, J=8.2  
Hz), 9.64 (br s, 1H).

Example 23

5 CIS- AND TRANS-2-BENZOYL-5-ACETOXY-1,3-OXATHIOLANE

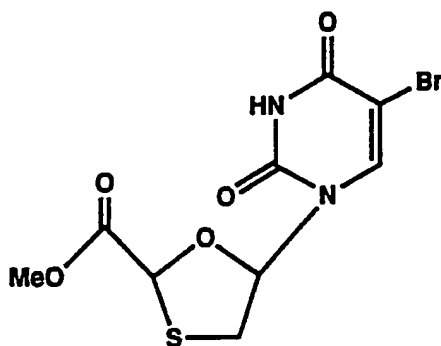


Phenyl glyoxal monohydrate (608 mg, 4.0 mmol)  
and 2,5-dihydroxy-1,4-dithiane (304 mg, 2.0 mmol) were  
heated for ca. 5 minutes at 65°C until the reagents  
melted. The reaction mixture was diluted with  
10 dichloromethane (40 mL). Pyridine (1.32 mL, 16.0  
mmol), 4-dimethylamino-pyridine (DMAP) (48 mg), and  
acetyl chloride (0.85 mL, 12.0 mmol) were added to the  
stirred solution at 0°C. The reaction mixture was  
stirred at room temperature for 4.5 hours and diluted  
15 with brine solution (15 mL). The organic layer was  
separated, washed with sodium bicarbonate and brine  
solutions, dried (sodium sulfate), and evaporated to a  
brown liquid (1.80 g). The residue was purified by  
silica gel chromatography eluting with hexanes:EtOAc  
20 (3:1) to yield the trans and cis isomers (2.4:1 ratio)  
(714 mg, 71%); <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 2.0 (s, 3H), 2.14 (s,  
3H), 3.15-3.25 (m, 1H), 3.35-3.45 (m, 1H), 6.42 (s,  
1H), 6.51 (s, 1H), 6.7 (m, 1H), 6.9 (m, 1H), 7.4-7.5  
(m, 2H), 7.55-7.65 (m, 1H), 7.9-8.0 (m, 2H).

Example 24CIS-2-(1'-PYRROLIDINOCARBONYL)-5-ACETOXY-1,3-OXATHIOLANE

To a solution of 5-acetoxy-oxathiolane-2-  
5 carboxylic acid (576 mg, 3.0 mmol), pyridine (0.533 mL,  
6.60 mmol), and dichloromethane (20 mL) at 0°C, was  
added oxalyl chloride (0.314 mL, 3.6 mmol). The  
reaction was stirred at 0°C for 30 minutes and then  
cooled to -70°C at which time pyrrolidine (0.5 mL, 6.0  
10 mmol) was added in one portion. The reaction was  
stirred at room temperature for 2 hours followed by  
addition of 1N HCl (5 mL). The organic layer was  
separated, washed with sodium bicarbonate and brine  
solutions, dried (sodium sulfate), and concentrated to  
15 yield 0.851 g of crude product. This residue was  
purified by silica gel chromatography eluting with  
EtOAc:hexanes (9:1) to give 616 mg (84%) of the desired  
product; <sup>1</sup>NMR (CDCl<sub>3</sub>) δ 1.80-2.00 (m, 4H), 2.11 (s, 3H),  
3.20-3.35 (m, 2H), 3.40-3.55 (m, 4H), 5.76 (s, 1H),  
20 6.60 (m, 1H).

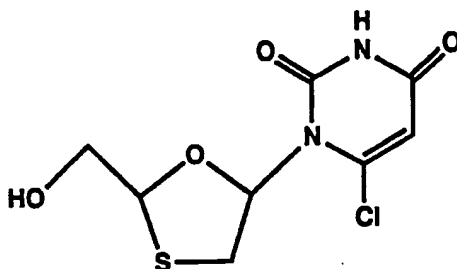
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Example 25CIS-2-CARBOMETHOXY-5-(5'-BROMOURACIL-1'-YL)-1,3-OXATHIOLANE

Bis-trimethylsilyl-acetamide (4 mL, 16.2  
5 mmol) was added to a suspension of 5-bromouracil (1.5  
g, 7.9 mmol) in dichloromethane (10 mL). The reaction  
was stirred for 30 minutes, yielding a clear solution.  
Then a dichloromethane solution (5 mL) of 2-  
carbomethoxy-5-acetoxy-1,3-oxathiolane (1.6 g, 7.8 mmol  
10 *cis:trans* 1:2) was added, followed by TMSI (1.1 mL, 7.7  
mmol).

The reaction was stirred at ambient  
temperature for 18 hours and then sequentially treated  
with saturated aqueous solutions of  $\text{Na}_2\text{S}_2\text{O}_3$  and  $\text{NaHCO}_3$   
15 to give a white suspension. The suspension was  
filtered to remove the solid (unreacted base). The  
filtrate was concentrated and triturated with EtOAc-  
Hex (1:1) to give white solid which was filtered,  
washed and dried to give 0.98 g (38%) of the product.  
20  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  3.2 (dd, 1H,  $J=7$  and 12 Hz), 3.47 (dd,  
1H,  $J=5$  and 12 Hz), 3.87 (s, 1H), 5.50 (s, 1H), 6.42  
(dd, 1H,  $J=5$  and 7 Hz), 8.72 (s, 1H), 9.19 (br s, 1H).

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Example 26CIS-2-HYDROXYMETHYL-5-(6'-CHLOROURACIL-1'-YL)-1,3-OXATHIOLANE

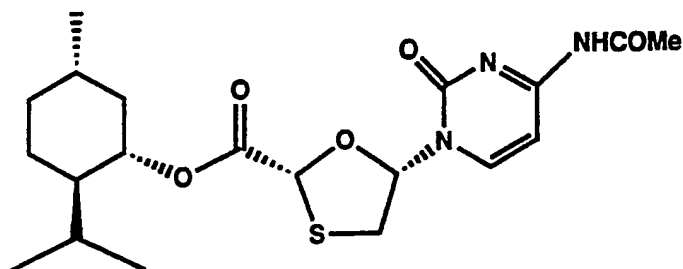
TMSOTf (4.5 mL, 27.3 mmol) was added to a  
5 suspension of bis-O-silyl-6-chlorouracil (9.5 g, 32.6  
mmol) and 2-carbethoxy-5-acetoxoxathiolane (6.3 g,  
27.4 mmol) in 1,2-dichloroethane (40 mL). The  
resulting clear solution was heated slowly up to 60°C  
and kept at this temperature for 1 hour, during which  
10 time a thick precipitate appeared. The reaction was  
cooled down to ambient temperature and the white  
precipitate was collected after filtration, washed and  
dried to give 3.5 g (42%) of the only *cis* nucleoside  
ester product (<sup>1</sup>H NMR). To a tetrahydrofuran (THF) (50  
15 mL) suspension of nucleoside ester product (2.6 g, 8.5  
mmol), under argon atmosphere, was slowly added LiBH<sub>4</sub>  
(0.4 g, 18.6 mmol). The reaction was stirred for 5  
hours, then quenched with methanol. The solvent was  
removed, followed by subjecting the resulting gummy  
20 material to column chromatography (2:2:1, EtOAc-Hex-  
MeOH, v/v) to yield 1.9 g (85%) of the title  
nucleoside. The overall yield of these two  
transformations was 64%; HPLC purity (96%); mp 202-  
204°C; <sup>1</sup>H NMR (DMSO-d<sub>6</sub>) δ 3.09-3.30 (1H), 3.38-3.47

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(1H), 3.60-3.72 (2H), 4.45 (1H), 5.05-5.09 (1H), 5.27 (1H), 5.59-5.62 (1H), 6.71-6.76 (1H);  $^{13}\text{C}$  NMR (DMSO- $d_6$ )  $\delta$  32.6, 63.2, 64.2, 84.7, 87.9, 94.4, 106.6, 128.6, 164.4.

5 Example 27

(1'S,2'R,5'S)-MENTHYL-5S-(N-4"-ACETYLCYTOSIN-1"-YL)-1,3-OXATHIOLANE-2R-CARBOXYLATE



To a stirred suspension of N-4-acetylcytosine (68 mg, 0.4 mmol) in dichloromethane (0.5 mL) containing 2,4,6-collidine (105  $\mu\text{L}$ , 0.8 mmol) under an argon atmosphere was added trimethylsilyl trifluoromethane-sulphonate (155  $\mu\text{L}$ , 0.8 mmol). The resulting mixture was stirred for 15 minutes to give a homogeneous solution. The substrate, (1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolane-2R-carboxylate (110 mg, 0.333 mmol) was introduced into the above solution in one batch. In a separate flask equipped with a condensor, a solution of hexamethyldisilazane (34  $\mu\text{L}$ , 0.167 mmol) and iodine (42 mg, 0.167 mmol) in dichloromethane (0.5 mL) was refluxed under argon atmosphere for 30 minutes. After it had cooled to room temperature, the purple solution formed was transferred, via a syringe, into the mixture containing the substrate and silylated base.

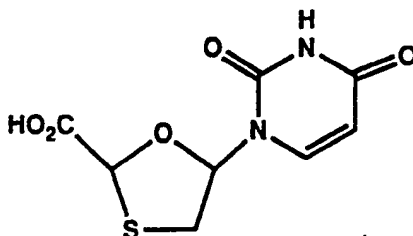


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The reaction mixture was kept at room temperature for 7 hours and then was quenched with a solution of a 1:1 mixture of saturated  $\text{NaHCO}_3$  and  $\text{Na}_2\text{S}_2\text{O}_3$ . The resulting mixture was stirred for 5 minutes and then was transferred to a separatory funnel with the aid of more dichloromethane. The aqueous phase was removed and the organic layer was washed with saturated  $\text{Na}_2\text{S}_2\text{O}_3$ , water, brine and then was dried ( $\text{Na}_2\text{SO}_4$ ). The solvent was removed under reduced pressure to provide 153 mg of crude product. To determine the ratio of the *cis*-[(1'S,2'R,5'S)-menthyl-5S-(N-4"-acetylcytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate] and *trans*-[(1'S,2'R,5'S)-menthyl-5R-(N-4"-acetylcytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate] product isomers, the crude product was subjected to  $^1\text{H}$  NMR analysis in  $\text{CDCl}_3$ . Judging from the signals of the C6 protons of the cytosine moiety, the ratio of *cis* [ $\delta$  8.70 (d,  $J=7.6$  Hz)] to *trans* [ $\delta$  7.79 (d,  $J=7.6$  Hz)] was determined to be 7:1.

20 Example 28

CIS-2-CARBOXYL-5-(URACIL-1'-YL)-1,3-OXATHIOLANE

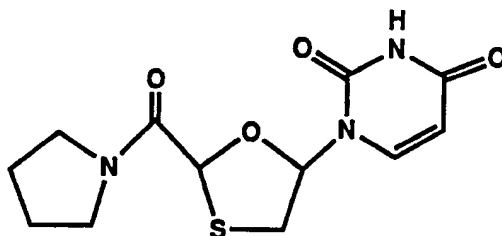


Iodotrimethylsilane (118  $\mu\text{L}$ , 0.832 mmol) was added to a stirred suspension of bis-trimethylsilyl-uracil (122 mg, 0.475 mmol) and *trans*-2-carboxyl-5-

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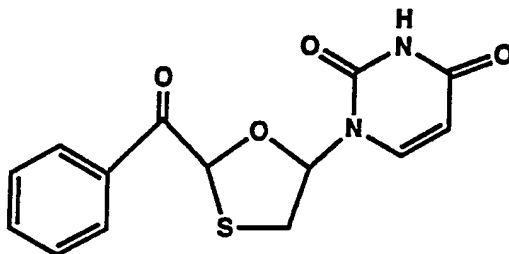
acetoxy-1,3-oxathiolane (76 mg, 0.396 mmol) in dichloromethane (2.5 mL) containing collidine (53  $\mu$ L, 0.396 mmol). The resultant mixture was stirred for 18 hours at room temperature under argon atmosphere and  
5 then was quenched by the addition of 5 mL of a 0.5 M solution of sodium carbonate. The aqueous phase was acidified with 1 M HCl solution to pH 4, followed by extraction with tetrahydrofuran (3x6 mL). The combined extract was dried over  $\text{MgSO}_4$  and the solvent was  
10 removed under reduced pressure. The crude product obtained was triturated with dichloromethane to provide a white suspension. The white solid was isolated by centrifugation and was dried under vacuum to afford 27 mg of the desired product whose  $^1\text{H}$  NMR spectrum  
15 indicated the presence of a small amount of uracil (ca. 10%) and an isomeric purity of  $\geq 95\%$ . The title compound displayed the following spectral characteristics:  $^1\text{H}$  NMR ( $\text{DMSO } d_6$ )  $\delta$  : 2.26 (d of d, 1H,  $J=4.9, 12.3$  Hz), 3.49 (d of d, 1H,  $J=5.2, 12.4$  Hz),  
20 5.57 (s, 1H), 5.71 (d of d, 1H,  $J=2.2, 8.0$  Hz; this signal collapsed to a doublet on treatment with  $\text{D}_2\text{O}$  ( $J=8.2$  Hz)), 6.29 (t, 1H,  $J=5.2$  Hz), 8.07 (d, 1H,  $J=8.2$  Hz), 11.41 (br s, 1H, exchanged with  $\text{D}_2\text{O}$  ).

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Example 29CIS 2-(1'-PYRROLIDINOCARBONYL)-5-(URACIL-1'-YL)-1,3-OXATHIOLANE

Iodotrimethylsilane (37  $\mu$ L, 1 equivalent) was  
5 added to a stirred solution of *cis* 2-(1'-pyrrolidino-  
carbonyl)-5-acetoxy-1,3-oxathiolane (64 mg, 0.26 mmol)  
and bis-trimethylsilyluracil (80 mg, 1.2 equivalents)  
in dichloromethane (1.5 mL) under argon atmosphere.  
The reaction mixture was kept for 1 hour and 20 minutes  
10 at room temperature. The reaction was quenched with a  
solution of a 1:1 mixture of saturated  $\text{Na}_2\text{S}_2\text{O}_3$  and  
 $\text{NaHCO}_3$  (2 mL), followed by dilution with  
dichloromethane (4 mL). The resultant mixture was  
stirred for 5 minutes and then was transferred to a  
15 separatory funnel with the aid of more dichloromethane.  
The aqueous phase was removed and the organic phase was  
washed with water, brine, and dried over anhydrous  
 $\text{Na}_2\text{SO}_4$ . Removal of the solvent under reduced pressure  
and subjection of the crude product thus obtained to  
20 column chromatography (7% MeOH-EtOAc) afforded the 74  
mg (95%) of the title compound;  $^1\text{H}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  1.85-  
2.00 (m, 2H), 2.00-2.15 (m, 2H), 3.25-3.70 (m, 6H), 5.61  
(s, 1H), 5.80 (d of d, 1H,  $J=2.3$ , 8.2 Hz), 6.44 (d of  
d, 1H,  $J=4.8$ , 7.0 Hz), 8.29 (br s, 1H), 8.88 (d, 1H,  
25  $J=8.1$  Hz).

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Example 30CIS 2-BENZOYL-5-(URACIL-1'-YL)-1,3-OXATHIOLANE

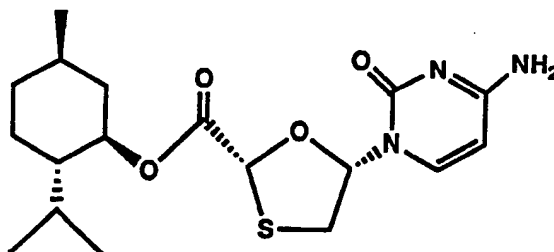
Trimethylsilyl trifluoromethanesulphonate (92  $\mu\text{L}$ , 0.475 mmol) was introduced to a stirred suspension of uracil (50 mg, 0.238 mmol) in dichloromethane (1.5 mL) containing collidine (63  $\mu\text{L}$ , 0.475 mmol) under argon atmosphere. The resultant mixture was stirred for 15 minutes to provide a homogeneous solution. A mixture (2.4:1, trans:cis) of 2-benzoyl-5-acetoxy-1,3-oxathiolane (50 mg, 0.198 mmol) was added as a solution in dichloromethane (1.5 mL), followed by iodotrimethylsilane (28  $\mu\text{L}$ , 0.198 mmol). The reaction was allowed to proceed for 22 hours and then was quenched with a solution of a 1:1 mixture of saturated  $\text{NaHCO}_3$  and  $\text{Na}_2\text{S}_2\text{O}_3$ . The resulting mixture was stirred for 5 minutes and then was transferred to a separatory funnel with the aid of more dichloromethane. The aqueous phase was removed and the organic layer was washed with saturated  $\text{Na}_2\text{S}_2\text{O}_3$ , water, brine and then was dried ( $\text{Na}_2\text{SO}_4$ ). Thin layer chromatography analysis of the crude product indicated that small amount of the starting material remain unreacted. The crude product was triturated with EtOAc to provide 26 mg (43%) of the

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title compound as a white solid;  $^1\text{H}$  NMR (DMSO):  $\delta$  3.19 (d of d, 1H, d of d,  $J=6.8, 12.1$  Hz), 3.60 (d of d, 1H,  $J=5.1, 12.2$  Hz), 5.77 (d, 1H,  $J=8.2$  Hz), 6.38 (d of d, 1H,  $J=5.2, 6.9$  Hz), 6.81 (s, 1H), 7.52-7.64 (m, 2H),  
5 7.66-7.76 (m, 1H), 7.94-8.04 (m, 2H), 8.22 (d, 1H,  $J=8.1$  Hz), 11.44 (br s, 1H).

Example 31

(1'R,2'S,5'R)-MENTHYL-5S-(CYTOSIN-1"-YL)-1,3-OXATHIOLANE-2R-CARBOXYLATE



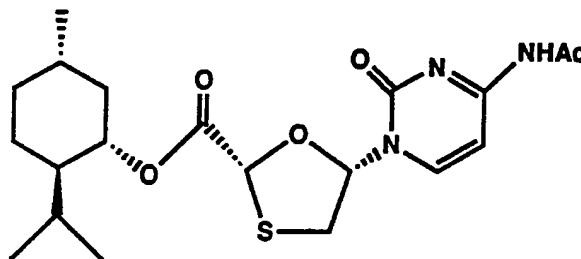
10 A 12:1 mixture of (1'R,2'S,5'R)-menthyl 5S-(N-4''-acetylcytosin-1''-yl)-oxathiolane-2R-oxathiolane carboxylate (*cis* isomer) and (1'R,2'S,5'R)-menthyl 5R-(N-4''-acetylcytosin-1''-yl)-oxathiolane-2R-oxathiolane carboxylate (*trans* isomer) (47 mg, 0.11 mmol) was  
15 dissolved in dichloromethane (0.5 mL) and 2-propanol (1 mL). Trifluoroacetic acid (0.2 mL) was added to this solution and the resultant mixture was heated at 60°C for 2 hours and then was kept at room temperature for 14.5 hours. The reaction mixture was diluted with  
20 dichloromethane and washed with saturated  $\text{NaHCO}_3$  solution, water, brine, and then was dried (anhydrous  $\text{Na}_2\text{SO}_4$ ). The solvent was removed under reduced pressure and the product obtained was dried under vacuum to

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afford 40 mg (95%) of the title compounds. The  $^1\text{H}$  NMR spectrum of the above material suggested a purity of  $\geq 97\%$ . Based on the signals derived from the C6 hydrogen of the cytosine moiety present in both of the isomers, the 12:1 ratio of the *cis* [ $\delta$  8.38 (d,  $J=7.3$  Hz)] and *trans* [ $\delta$  7.48 (d,  $J=7.3$  Hz)] nucleosides was maintained. The major compound was obtained by fractional crystallization with methanol and displayed physical properties identical to those reported in this example.

#### Example 32

(1'S,2'R,5'S)-MENTHYL-5S-(N-4"-ACETILCYTOSIN-1"-YL)-1,3-OXATHIOLANE-2R-CARBOXYLATE



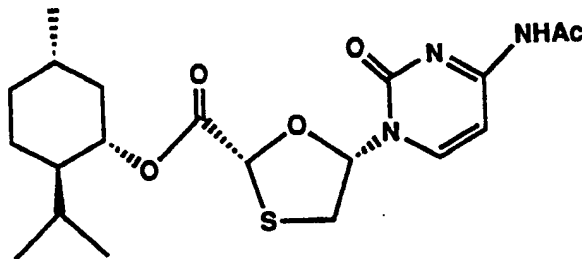
(1'S,2'R,5'S)-menthyl 5R-acetoxy-1,3-oxathiolane-2R-carboxylate (55 mg, 0.166 mmol) in dichloromethane (0.5 mL) and iodotrimethylsilane (0.026 mL, 0.166 mmol) were added to monosilylated N-4-acetylcytosine (59 mg, 0.198 mmol), generated by refluxing N-4-acetylcytosine in 1,1,1,3,3,3-hexamethyl-20 disilazane (HMDS) overnight in the presence of catalytic amount of ammonium sulfate and subsequently removing HMDS, in dichloromethane (0.5 mL) under argon atmosphere at room temperature. The stirring was continued for 19 hours and thin layer chromatography 25 showed almost complete consumption of the starting

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oxathiolane. The reaction mixture was diluted with dichloromethane, washed with saturated aqueous sodium bicarbonate, aqueous sodium thiosulfate and brine, dried over sodium sulfate, concentrated and dried to afford 70 mg (100%) of crude products. <sup>1</sup>H NMR suggested *cis:trans* ratio at 15:1 and the presence of ca. 4.6% of unreacted oxathiolane. <sup>1</sup>H NMR (CDCl<sub>3</sub>): 0.78 (d, 3H), 0.80-2.10 (m, 15H), 2.27 (s, 3H), 3.12-3.30 (m, 1H), 3.52-3.78 (m, 1H), 4.78 (m, 1H), 5.51 (s, 0.896H), 5.60 (s, 0.046H), 5.82 (s, 0.058H), δ 6.42 (t, 0.896H), 6.63 (dd, 0.046 H), 6.68 (d, 0.058H), 7.47 (d, 0.954H), 7.77 (d, 0.058H), 8.70 (d, 0.896H). The major compound was isolated by crystallization from methanol or trituration with ethylacetate-ether mixtures.

15 Example 33

(1'S,2'R,5'S)-MENTHYL-5S-(N-4"-ACETYLCYTOSIN-1"-YL)-1,3-OXATHIOLANE-2R-CARBOXYLATE



2,6-lutidine (0.023 mL, 0.199 mmol) and trimethylsilyl trifluoromethanesulfonate (0.038 mmol, 0.199 mmol) were added to N-4-acetylcytosine (30.5 mg, 0.199 mmol) in dichloromethane (0.2 mL) at room temperature under argon atmosphere. The mixture was stirred for 20 minutes and a solution of (1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolane-2R-carboxylate (55 mg, 0.166 mmol) in dichloromethane (0.3 mL) and iodotrimethyl-silane (0.026 mL, 0.166 mmol) were

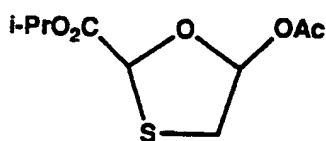
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introduced successively. The stirring was continued for 2.5 hour and thin layer chromatography showed complete consumption of the starting oxathiolane. The reaction mixture was diluted with dichloromethane, washed with saturated aqueous sodium bicarbonate, aqueous sodium thiosulfate and brine, dried over sodium sulfate, concentrate and dried to afford 70 mg (100%) of crude products. <sup>1</sup>H NMR suggested *cis:trans* ratio at 10:1 and no other impurity detectable by the spectrum.

<sup>1</sup>H NMR (CDCl<sub>3</sub>): 0.78 (d, 3H), 0.80-2.10 (m, 15H), 2.27 (s, 3H), 3.16 (dd, 0.91H), 3.25 (d, 0.09H), 3.63 (dd, 0.91H), 3.74 (dd, 0.09H), 4.78 (m, 1H), 5.51 (s, 0.91H), 5.82 (s, 0.09H); δ 6.42 (t, 0.91H), 6.68 (d, 0.09H), 7.47 (d, 1H), 7.77 (d, 0.09H), 8.70 (d, 0.91H).

#### Example 34

#### CIS- AND TRANS- ISOPROPYL 5-ACETOXY-1,3-OXATHIOLANE-2-CARBOXYLATE



A solution of *cis-* and *trans* 5-acetoxy-1,3-oxathiolane-2-carboxylic acid (260 mg, 1.3528 mmol) and isopropanol (0.11 mL, 1.3528 mmol) in dichloromethane (4 mL) at 0°C was treated with dicyclohexylcarbodiimide (DCC) (279 mg, 1.3528 mmol) in dichloromethane (1 mL) and 4 dimethylaminopyridine (DMAP) (14 mg, 0.135 mmol).

The mixture was stirred at room temperature overnight,



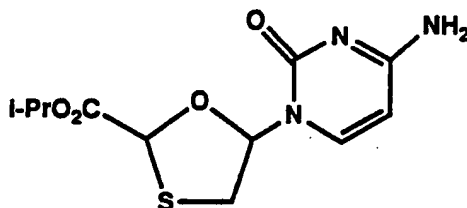
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then diluted with ether and filtered through a Celite® pad. The filtrate was concentrated and the residue was chromatographed on silica gel with ethyl acetate-hexane to give the products as a colorless oil (263 mg, 83%).

- 5 <sup>1</sup>H NMR (CDCl<sub>3</sub>): δ 1.26 (6H, d); 2.10, 2.11 (3H, s); 3.13-3.46 (2H, m); 5.05 (1H, m); 5.60, 5.61 (1H, s); 6.63 (0.54H, m); 6.78 (0.46H, d).

Example 35

- 10 CIS-ISOPROPYL-5-(CYTOSIN-1'-YL)-1,3-OXATHIOLANE-2-CARBOXYLATE



- 2,4,6-collidine (0.23 mL, 1.74 mmol) and t-butyl-dimethylsilyl trifluoromethanesulfonate (0.4 mL, 1.74 mmol) were added to a suspension of cytosine (96.7 mg, 0.87 mmol) in dichloromethane (0.8 mL) at room temperature under argon atmosphere. The mixture was stirred for 25 minutes and a solution of *cis:trans* (1.2:1) isopropyl 5-acetoxy-1,3-oxathiolane-2-carboxylate (168 mg, 0.717 mmol) in dichloromethane (0.8 mL) and a solution of iodotrimethylsilane (0.114 mL, 0.788 mmol) were introduced successively. Stirring was continued for one hour and the reaction mixture was diluted with dichloromethane, washed with saturated aqueous sodium thiosulfate, water and brine, dried over sodium sulfate and concentrated. The residue was trituated with ether-hexane (1:1, 7 mL) and saturated
- 15  
20  
25

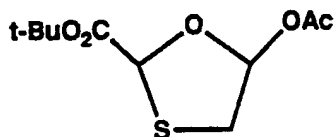
- 80 -

aqueous sodium bicarbonate (1.5 mL). The aqueous layer was removed and the remaining mixture was centrifuged.

The solid was washed twice with hexanes and the washings were combined with centrifugate, washed  
5 with 1N HCl, water and brine, dried and concentrated to give the unreacted starting material in virtually pure form (64 mg, 38%, *cis:trans*=1:9). The white solid was dried and gave the products as a *cis:trans* mixture in  
12:1 ratio (122.6 mg, 60%). <sup>1</sup>H NMR (CDCl<sub>3</sub>): δ 1.30 (t, 6H), 3.11 (dd, 1H), 3.52 (dd, 1H), 5.11 (m, 1H), 5.45 (s, 1H), 5.82 (d, 1H), 6.47 (dd, 0.92H), 6.72 (m, 0.08H), 7.49 (d, 0.08H), 8.32 (d, 0.92H).

#### Example 36

15 CIS- AND TRANS- T-BUTYL 5-ACETOXY-1,3-OXATHIOLANE-2-CARBOXYLATE



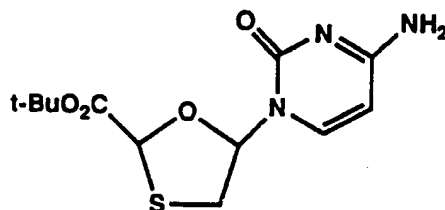
A solution of *cis-* and *trans-* 5-acetoxy-1,3-oxathiolane-2-carboxylic acid (176 mg, 0.915 mmol) and *t*-butanol (0.095 mL, 0.915 mmol) in dichloromethane (4 mL) at 0°C was treated with DCC (207 mg, 1 mmol) in  
20 dichloromethane (1 mL) and DMAP (11 mg, 0.09 mmol). The mixture was stirred at room temperature overnight, then diluted with ether and filtered through a Celite® pad. The filtrate was concentrated and the residue was chromatographed on silica gel with ethyl acetate-hexane  
25 to give the products as a colorless oil (175 mg, 77%).

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$^1\text{H}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  1.46 (9H, d); 2.07, 2.09 (3H, s); 3.10-3.44 (2H, m); 5.50, 5.52 (1H, s); 6.60 (0.42H, m); 6.74 (0.58H, d).

Example 37

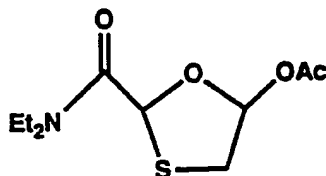
5 CIS-T-BUTYL-5-(CYTOSIN-1'-YL)-1,3-OXATHIOLANE-2-CARBOXYLATE



2,4,6-collidine (0.187 mL, 1.4 mmol) and t-butyl-dimethylsilyl trifluoromethanesulfonate (0.325 mL, 1.4 mmol) were added to a suspension of cytosine  
10 (78.6 mg, 0.7 mmol) in dichloro-methane (0.6 mL) at room temperature under argon atmosphere. The mixture was stirred for 25 minutes and a mixture of cis and trans (1:1.4) t-butyl 5-acetoxy-1,3-oxathiolane-2-carboxylate (146.5 mg, 0.59 mmol) in dichloromethane  
15 (0.6 mL) and iodotrimethylsilane (0.092 mL, 0.65 mmol) were introduced successively. Stirring was continued for one hour and the reaction mixture was diluted with dichloromethane, washed with saturated aqueous sodium thiosulfate, water and brine, dried over sodium sulfate  
20 and concentrated. The residue was triturated with ether-hexanes (1:1, 7 mL) and saturated aqueous sodium bicarbonate (1.5 mL). The aqueous layer was removed and the remaining mixture was centrifuged. The solid was washed twice with hexanes and the washings were

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combined with the centrifugate, washed with 1N HCl, water and brine, dried and concentrated to give the unreacted starting material in virtually pure form (77 mg, 52.6%, *cis:trans*=1:11). The white solid was dried  
5 and gave the products as a *cis:trans* mixture in 16:1 ratio (82.6 mg, 46.4%). <sup>1</sup>H NMR (CDCl<sub>3</sub>): δ 1.50, 1.52 (s, 9H), 3.12 (dd, 0.94H), 3.20 (dd, 0.06H), 3.52 (dd, 0.94H), 3.72 (dd, 0.06H), 5.37 (s, 0.94H), 5.75 (s, 0.06H), 5.82 (d, 1H), 6.44 (dd, 0.94H), 6.71 (d,  
10 0.06H), 7.49 (d, 0.06H), 8.38 (d, 0.98H).

Example 38CIS- AND TRANS-2-N,N-DIETHYLAMINOCARBONYL-5-ACETOXY-1,3-OXATHIOLANE

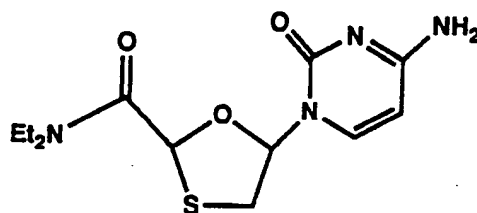
A solution of *cis*- and *trans*-5-acetoxy-1,3-oxathiolane-2-carboxylic acid (119 mg, 0.62 mmol) and diethylamine (0.07 mL, 0.68 mmol) in dichloromethane (2 mL) at 0°C was treated with DCC (140 mg, 0.68 mmol) in dichloromethane (1 mL) and DMAP (7.6 mg, 0.06 mmol). The mixture was stirred at room temperature overnight,  
20 then diluted with ether and filtered through a Celite® pad. The filtrate was concentrated and the residue was chromatographed on silica gel with ethyl acetate-hexane to give the products as a colorless oil (84.5 mg, 55%).  
<sup>1</sup>H NMR (CDCl<sub>3</sub>): δ 1.10, 1.40 (6H, t); 2.07, 2.10 (3H,

- 83 -

s); 3.15-3.56 (6H, m); 5.80, 5.87 (1H, s); 6.58 (0.53H, m); 6.83 (0.47H, d).

Example 39

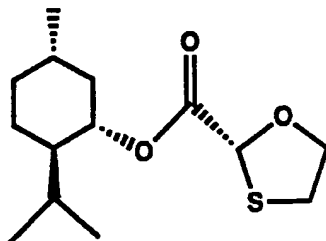
5 CIS-2-N,N-DIETHYLAMINOCARBONYL-5-(CYTOSIN-1'-YL)-1,3-OXATHIOLANE



2,4,6-collidine (0.108 mL, 0.82 mmol) and t-butyl-dimethylsilyl trifluoromethanesulfonate (0.188 mL, 0.82 mmol) were added to a suspension of cytosine (45.5 mg, 0.41 mmol) in dichloromethane (0.4 mL) at  
10 room temperature under argon atmosphere. The mixture was stirred for 25 minutes and a mixture of *cis* and *trans* (1.12:1) 2-N,N-diethylaminocarbonyl-5-acetoxy-1,3-oxathiolane (84 mg, 0.34 mmol) in dichloromethane (0.4 mL) and a solution of iodotrimethylsilane (0.053  
15 mL, 0.375 mmol) were introduced successively. Stirring was continued for one hour and the reaction mixture was diluted with dichloromethane, washed with saturated aqueous sodium thiosulfate, water and brine, dried over sodium sulfate and concentrated. The residue was  
20 triturated with ether-hexane (1:1, 7 mL) and saturated aqueous sodium bicarbonate (1.5 mL). The aqueous layer was removed and the remaining mixture was centrifuged. The solid was washed twice with hexanes and the washings were combined with centrifugate, washed with  
25 1N HCl, water and brine, dried and concentrated to give

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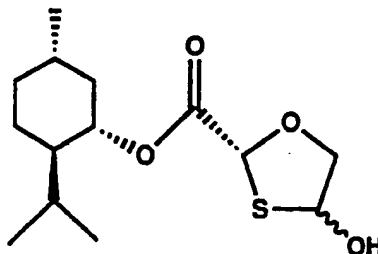
the unreacted starting material in virtually pure form (17 mg, 20%, trans only). The white solid was dried to give the products as a *cis:trans* mixture in 24:1 ratio (47.5 mg, 47.5%).  $^1\text{H}$  NMR ( $\text{DMSO}-d_6$ ):  $\delta$  1.04 (t, 3H,  $J=7$  Hz), 1.12 (t, 3H,  $J=7$  Hz), 3.17 (dd, 1H,  $J=5$  Hz, 9 Hz), 3.30 (m, 4H), 3.53 (dd, 1H,  $J=5$  Hz, 9 Hz), 5.74 (d, 1H,  $J=7$  Hz), 5.96 (s, 1H), 6.28 (t, 0.96H,  $J=5$  Hz), 6.62 (m, 0.04H), 7.16 (b.s., NH), 7.22 (b.s., NH), 7.60 (d, 0.04H), 8.46 (d, 0.96H,  $J=7$  Hz).

10 Example 40(1'S,2'R,5'S)-MENTHYL-1,3-OXATHIOLANE-2R-CARBOXYLATE

To a mixture of (1'S,2'R,5'S)-menthyl-5R-acetoxy-1,3-oxathiolane-2R-carboxylate (2.01 g, 6.08 mmol) and triethylsilane (9.67 mL, 60.05 mmol) at room temperature under argon atmosphere was added trimethylsilyl trifluoromethanesulfonate (1.17 mL, 6.04 mmol). The reaction mixture was stirred at room temperature for 12 hours, then diluted with dichloromethane, washed with saturated aqueous solution of sodium bicarbonate, dried over anhydrous sodium sulfate, and evaporated to dryness *in vacuo* to afford crude product. Subsequent chromatography on silica gel using hexane-ethyl acetate as eluate gave the product as colourless oil (1.33 g, 80.5%)  $^1\text{H}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  0.75-2.10 (m, 15H), 2.97-3.20 (m, 2H), 4.20-4.40 (m, 2H), 4.72 (dt, 1H), 5.45 (s, 1H)  $[\alpha]_D^{+104^\circ}$  (c 1.16,  $\text{CHCl}_3$ ).

Example 41

(1'S,2'R,5'S)-MENTHYL-4R-HYDROXY-1,3-OXATHIOLANE-2R-CARBOXYLATE AND (1'S,2'R,5'S)-MENTHYL-4S-HYDROXY-1,3-OXATHIOLANE-2R-CARBOXYLATE



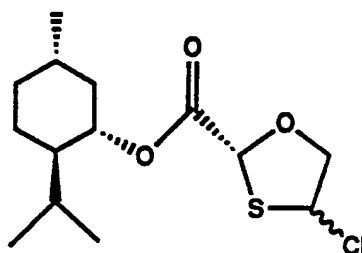
5           A mixture of (1'S,2'R,5'S)-menthyl-1,3-oxathiolane-2R-carboxylate (0.500 g, 1.84 mmol) and benzoylperoxide (0.489 g, 97%, 1.96 mmol) in 20 mL benzene was heated to reflux for 6 hours. The organic solvent was removed *in vacuo* and the residue was  
10 diluted with dichloromethane, washed with saturated aqueous solution of sodium bicarbonate, dried over anhydrous sodium sulfate, and evaporated to dryness *in vacuo* to afford crude benzoate product. Subsequent chromatography by using hexane-ethyl acetate as eluate  
15 gave the benzoate as a solid (0.21 g, 30.3%). The mixture of the benzoate (0.200 g, 0.531 mmol) and potassium carbonate (0.073 g, 0.532 mmol) in THF-MeOH-H<sub>2</sub>O (4 mL/5 mL/2 mL) was stirred at 0°C for 7 hours and organic solvent was removed *in vacuo*. The residue was  
20 diluted with H<sub>2</sub>O (7 mL), extracted with ether (10 mL), acidified with aqueous HCl, and extracted with dichloromethane. The dichloromethane layer was dried over sodium sulfate and evaporated to dryness *in vacuo* to afford crude product. Subsequent chromatography  
25 using hexane ether as eluent gave the product as a

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solid (67 mg, 43.7%)  $^1\text{H}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  0.75-2.10 (m, 15H), 4.03-4.83 (m, 2H), 5.52-5.75 (m, 2H).

Example 42

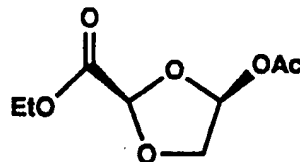
5 (1'S,2'R,5'S)-MENTHYL-4R-CHLORO-1,3-OXATHIOLANE-2R-CARBOXYLATE AND (1'S,2'R,5'S)-MENTHYL-4S-CHLORO-1,3-OXATHIOLANE-2R-CARBOXYLATE



To a mixture of (1'S,2'R,5'S)-menthyl-4R-hydroxy-1,3-oxathiolane-2R-carboxylate and (1'S,2'R,5'S)-menthyl-4S-hydroxy-1,3-oxathiolane-2R-carboxylate (40mg, 0.138 mmol) and methytrifluoromethansulfonyl chloride (18.24  $\mu\text{L}$ , 0.239 mmol) in dichloromethane (5 mL) at room temperature under argon atmosphere was added triethylamine (57.99 mL, 0.416 mmol). The reaction mixture was stirred at 15 room temperature for 2 hours then diluted with dichloromethane, washed with saturated aqueous solution of sodium bicarbonate, dried over anhydrous sodium sulfate, and evaporated to dryness *in vacuo* to afford crude product. Subsequent chromatography by using 20 hexane ether as eluent gave the product as two diastereomers (18 mg, 42.3%, 14.6 mg, 34.2%) epimeric at C4.  $^1\text{H}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  0.75-2.05 (m, 15H), 4.55 (m, 1H), 4.69 (m, 1H), 5.75 (m, 1H), 5.80 (m, 1H);  $\delta$  0.75-2.10 (m, 15H), 4.33 (m, 1H), 4.78 (m, 1H), 5.56 (s, 25 1H), 5.68 (m, 1H).

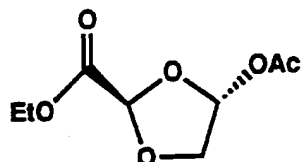


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Example 43CIS 2-CARBOETHOXY-4-ACETOXY-1,3-DIOXOLANE

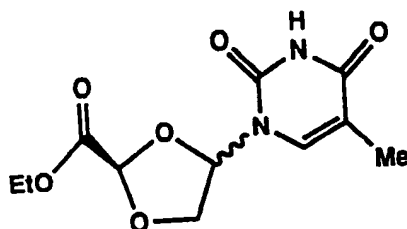
A 2.5:1 mixture of *cis* and *trans*-2-carboethoxy-4-acetyl-1,3-dioxolane (406 mg, 2.16 mmol),  
5 85% meta-chloroperbenzoic acid (mCPBA) (68 mg, 3.81 mmol) and sodium carbonate (389 mg, 3.67 mmol) in dry dichloromethane (10 mL) was stirred under argon for 16 hours at room temperature. The resultant suspension was diluted with dichloromethane and water and stirred  
10 for 10 minutes. The aqueous phase was removed and the organic phase was washed successively with saturated sodium thiosulfate, water, brine and then was dried over anhydrous magnesium sulfate. The solvent was removed under reduced pressure and the crude product  
15 thus obtained was subjected to flash column chromatography (30% EtOAc-Hexanes) to give the title compound (11% yield) which displayed the following spectral characteristics;  $^1\text{H}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  1.31 (t, 3H,  $J=7.2$  Hz), 2.07 (s, 1H), 4.15 (d of d, 1H,  $J=4.5$ , 9.1 Hz), 4.21-4.29 (m, 3H), 5.42 (s, 1H), 6.39 (d of d, 1H,  $J=2.4$ , 4.5 Hz);  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  14.05, 20.97, 29.69, 71.34, 94.04, 99.80, 167.19, 170.11.

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Example 44TRANS 2-CARBOETHOXY-4-ACETOXY-1,3-DIOXOLANE

A 2.5:1 mixture of *cis* and *trans*-2-carboethoxy-4-acetyl-1,3-dioxolane (406 mg, 2.16 mmol),  
5 85% mCPBA (68 mg, 3.81 mmol) and sodium carbonate (389 mg, 3.67 mmol) in dry dichloromethane (10 mL) was stirred under argon for 16 hours at room temperature. The resultant suspension was diluted with dichloromethane and water and stirred for 10 minutes.  
10 The aqueous phase was removed and the organic phase was washed successively with saturated sodium thiosulfate, water, brine and then was dried over anhydrous magnesium sulfate. The solvent was removed under reduced pressure and the crude product thus obtained  
15 was subjected to flash column chromatography (30% EtOAc-Hexanes) to give the title compound (49% yield) which displayed the following spectral characteristics;  
 $^1\text{H}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  1.29 (t, 3H,  $J=7.2$  Hz), 2.09 (s, 1H), 4.12 (d of d, 1H,  $J=0.9, 9.1$  Hz), 4.19-4.31 (m, 3H),  
20 5.53 (s, 1H), 6.48 (d of d, 1H,  $J=0.9, 3.9$  Hz).

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Example 45CIS AND TRANS 2-CARBOETHOXY-4-(THYMIN-1'-YL)-1,3-DIOXOLANE

To a stirred suspension of thymine (44.5 mg,  
5 0.353 mmol) in dichloromethane (1 mL) containing 2,6-lutidine (82  $\mu$ L, 0.706 mmol) under an argon atmosphere was added trimethylsilyl trifluoromethanesulphonate (136  $\mu$ L, 0.706 mmol). The resulting mixture was stirred for 15 minutes to give a homogeneous solution.

10 A solution of the substrate, ethyl 4-acetoxy-1,3-dioxolane-2-carboxylate (60 mg, 0.294 mmol) in dichloromethane (1 mL) and iodotrimethylsilane (42  $\mu$ L, 0.294 mmol) was sequentially introduced into the above solution. The reaction mixture was stirred at room

15 temperature for 5 hours and then was quenched with a half-saturated solution of  $\text{Na}_2\text{S}_2\text{O}_3$  (2 mL), followed by dilution with dichloromethane (5 mL). The resulting mixture was stirred for 5 minutes and then was transferred to a separatory funnel with the aid of more

20 dichloromethane. The aqueous phase was removed and the organic layer was washed with saturated  $\text{Na}_2\text{S}_2\text{O}_3$ , water, 1 M HCl, brine and then was dried ( $\text{Na}_2\text{SO}_4$ ). The solvent was removed under reduced pressure to provide the crude product. This material was suspended in

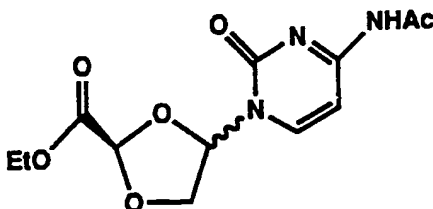
25 dichloromethane (~ 1.5 mL) and then was triturated with a 1:1 mixture of EtOAc-Hexane (~ 6 mL) to give 25 mg of

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the *cis* nucleoside as a white solid;  $^1\text{H}$  NMR ( $\text{DMSO } d_6$ ):  $\delta$  1.23 (t, 3H,  $J=7.1$  Hz), 1.78 (d, 3H,  $J=1$  Hz), 4.15-4.30 (m, 4H), 4.38 (d of d, 1H,  $J=2.3, 9.8$  Hz), 5.33 (s, 1H), 6.33 (d of d, 1H,  $J=2.3, 5.8$  Hz), 7.52 (d, 1H,  $J=1.1, 1\text{ Hz}$ ), 11.42 (br s, 1H). The triturate was concentrated and subjected to column chromatography (70% EtOAc-Hexane) to afford 26 mg of the two nucleoside as a 1:1 mixture;  $^1\text{H}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  1.33 (t, 1.5H,  $J=7.2$  Hz), 1.35 (t, 1.5H,  $J=7.2$  Hz), 1.91-1.99 (two overlapping d, 3H), 4.16 (d of d, 0.5H,  $J=1.9, 9.7$  Hz), 4.20-4.38 (m, 3H), 4.53 (d of d, 0.5H,  $J=5.8, 9.7$  Hz), 5.30 (s, 0.5H), 5.72 (s, 0.5H), 6.44 (d of d, 0.5H,  $J=3.3, 5.4$  Hz), 6.60 (d of d, 0.5H,  $J=2.0, 5.8$  Hz), 7.10 (d, 0.5H,  $J=1.3$  Hz), 7.75 (d, 0.5H,  $J=1.3$  Hz), 9.40 (br s, 0.5H), 9.43 (br s, 0.5H).

#### Example 46

#### CIS AND TRANS 2-CARBOETHOXY-4-(N-4'-ACETILCYTOSIN-1'-YL)-1,3-DIOXOLANE



To a stirred suspension of N-acetylcytosine  
. 20 (66 mg, 0.430 mmol) in dry  $\text{CH}_2\text{Cl}_2$  (1.5 mL) under an argon atmosphere was added, successively, 2,6-lutidine (100  $\mu\text{L}$ , 0.859 mmol) and trimethylsilyl trifluoromethanesulphonate (166  $\mu\text{L}$ , 0.859 mmol). The resultant mixture was stirred for 25 minutes to produce

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a homogeneous solution. A solution of a 4:1 mixture of *cis* and *trans*-2-carboethoxy-4-acetoxy-1,3-dioxolane (73 mg, 0.358 mmol) in  $\text{CH}_2\text{Cl}_2$  (1 mL) was then introduced, followed by iodotrimethylsilane (51  $\mu\text{L}$ , 0.358 mmol).

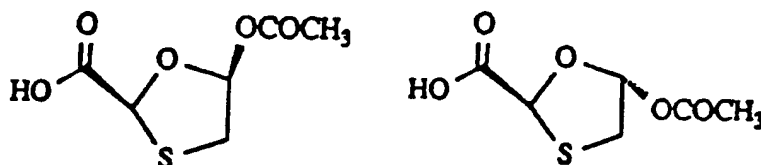
5 The reaction was allowed to proceed for 16 hours and then was quenched with saturated sodium thiosulfate. The resulting mixture was diluted with  $\text{CH}_2\text{Cl}_2$  and was washed successively with saturated sodium thiosulfate, water, brine, and then was dried over anhydrous

10 magnesium sulfate. Removal of the solvent under reduced pressure gave the crude product which was purified by flash column chromatography (2% MeOH-EtOAc) to afford 44% of the title compounds as a 3:1 mixture of the *cis* and *trans* isomers;  $^1\text{H}$  NMR ( $\text{CDCl}_3$ ):  $\delta$  1.34 (t, 3H,  $J=7.0$  Hz), 2.28 (s, 0.75H), 2.29 (s, 0.25H), 4.21-4.35 (m, 3H), 4.36 (d of d, 0.75 H,  $J=5.2$ , 9.9 Hz), 4.59 (d of d, 0.25H,  $J=5.2$ , 9.9 Hz), 5.39 (s, 0.75H), 5.77 (s, 0.25H), 6.24 (d of d, 0.75H,  $J=2.8$ , 5.1 Hz), 6.39 (d of d, 0.25H,  $J=1.7$ , 5.1 Hz), 7.49 (2

15 3H,  $J=7.0$  Hz), 2.28 (s, 0.75H), 2.29 (s, 0.25H), 4.21-4.35 (m, 3H), 4.36 (d of d, 0.75 H,  $J=5.2$ , 9.9 Hz), 4.59 (d of d, 0.25H,  $J=5.2$ , 9.9 Hz), 5.39 (s, 0.75H), 5.77 (s, 0.25H), 6.24 (d of d, 0.75H,  $J=2.8$ , 5.1 Hz), 6.39 (d of d, 0.25H,  $J=1.7$ , 5.1 Hz), 7.49 (2

20 overlapping doublets, 1H), 7.79 (d, 0.25H,  $J=7.6$  Hz), 8.40 (d, 0.75H,  $J=7.6$  Hz), 9.95 (br s, 1H).

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Example 47(±)-CIS AND TRANS-5-ACETOXY-1,3-OXATHIOLANE-2-CARBOXYLIC ACID

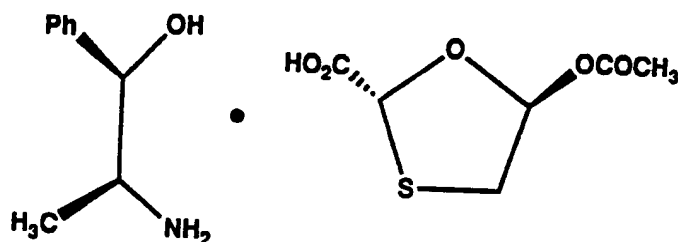
*Trans*-5-hydroxy-1,3-oxathiolane-2-carboxylic  
5 acid (250 g, 1.67 mol) was added, in portions, to a stirred solution of acetic anhydride (0.625 L, 6.62 mol) and methanesulphonic acid (5 mL, 77 mmol) at room temperature. The resultant clear solution was stirred at room temperature for 60 minutes, slowly added to  
10 stirred aqueous 0.03 M sodium bicarbonate solution (2.5L) and then the mixture was stirred for a further 60 minutes. Sodium chloride (750 g, 12.83 mol) was added and the mixture was stirred for a further 30 minutes, clarified, and then extracted with isopropyl  
15 acetate (1 x 1.25 L, 3 x 0.625 L). The combined extracts were concentrated to 1.25 L under reduced pressure. Xylene (2.5 L) was added and the mixture reconcentrated to 1.25 L under reduced pressure. The xylene addition/reconcentration procedure was repeated  
20 and the resultant suspension was cooled to room temperature and stirred for 18 hours. The solid was collected by vacuum filtration, washed with xylene (2 x 0.25 L) and dried, in vacuo, at 40-45° to give the title compound (265 g, 83%) which was shown, by

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comparison of  $^1\text{H}$  NMR spectra, to be a 65:35 mixture of the compounds of Examples 3 and 4.

Example 48

5 5R-ACETOXY-1,3-OXATHIOLANE-2R-CARBOXYLIC ACID, SALT WITH 1S,2R- $\alpha$ -(1-AMINOETHYL)BENZENEMETHANOL (1:1)



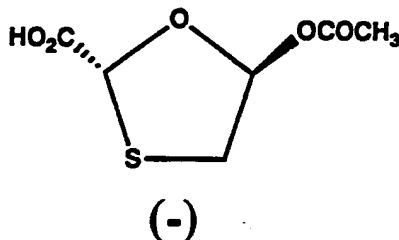
a) A solution of 1S,2R- $\alpha$ -(1-aminoethyl)-benzenemethanol (125.9 g, 0.83 mol) in isopropyl acetate (0.5L) was added to a stirred solution of ( $\pm$ ) *cis*-/trans-5-acetoxy-1,3-oxathiolane-2-carboxylic acid  
 10 (Example 47; 400 g, 2.08 mol), in isopropyl acetate (4.2 L), at room temperature under a nitrogen atmosphere. The resultant solution was stirred for 10 minutes, seeded with authentic product (0.4 g) and stirred for a further 4 hours at room temperature. The  
 15 suspension was stirred at 15-18° for 17 hours and the solid was collected by vacuum filtration, washed with isopropyl acetate (1 x. 0.4 L, 1 x 0.2 L) and dried, in *vacuo*, at 45° to give the title compound (205.9 g, 28%).  $[\alpha]_D^{+34}$  (MeOH), mp 151-2° (decomp),  $\delta$ (DMSO- $\text{D}_6$ )  
 20 0.91 (d, 3H, J=6.8 Hz), 2.05 (s, 3H), 3.04 (d, 1H, J=11 Hz), 3.32 (dd, 1H, J=4.2 Hz), 3.40 (dq, 1H, J=6.8, 2.4 Hz), 4.97 (d, 1H, J=2.4 Hz), 5.34 (s, 1H), ca. 6.4 (br, 1H), 7.2-7.4 (m, 5H), ca. 8.3 (br, 3H).

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b) A solution of 1S,2R- $\alpha$ -(1-aminoethyl)-benzenemethanol (177 mg, 1.17 mmol) in isopropyl acetate (1 mL) was added to a stirred solution of ( $\pm$ )-trans-5-acetoxy-1,3-oxathiolane-2-carboxylic acid (500 mg, 2.60 mmol) in isopropyl acetate (6 mL) at 25-30°, and further isopropyl acetate (0.5 mL) was added. Crystallisation commenced after 5 minutes. The suspension was stirred at 25-30° for 18 hours and then the solid was collected by vacuum filtration, washed with isopropyl acetate (1 mL) and dried, in vacuo, at 40° to give the title compound (353 mg, 40%), as shown by comparison of its  $^1\text{H}$  NMR spectrum with that of part (a).

Example 49

15 (-)-TRANS-5-ACETOXY-1,3-OXATHIOLANE-2-CARBOXYLIC ACID



5 M-Aqueous hydrochloric acid (126 mL, 0.63 mol) was added to a stirred suspension of the compound of Example 48 (180 g, 0.52 mol) in saturated aqueous sodium chloride (414 mL) at room temperature. The mixture was stirred at room temperature for 30 minutes, cooled to 10° and stirred at this temperature for a further 30 minutes. The solid was collected by vacuum filtration, washed with chilled water (2 x 90 mL) and

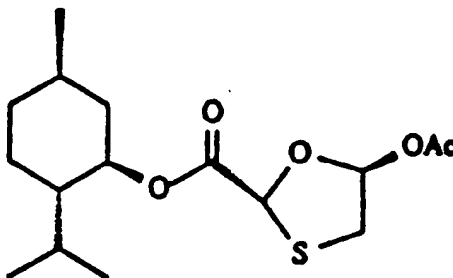


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dried, in *vacuo*, at 33° to give the title compound (81.3 g, 81%).

Example 50

5 (1'R,2'S,5'R)-MENTHYL-5R ACETOXY-1,3-OXATHIOLANE-2R-CARBOXYLATE



a) A solution of oxalyl chloride (66.5 g, 0.52 mol) in dichloromethane (120 mL) was added over 30 minutes to a stirred cold (-5°) mixture of N,N-dimethylformamide (32 mL) and dichloromethane (240 mL),  
10 and the suspension formed was stirred at -5 to 0° for 30 minutes. The compound of Example 49 (80 g, 0.42 mol) was added in portions and the resultant yellow solution was stirred at 0° for 45 minutes. This solution was added over 60 minutes to a stirred, cold  
15 (-5°) solution of (1R,2S,5R)-(-)-menthol (65.2 g, 0.425 mol) in dichloromethane (200 mL) and pyridine (84mL, 1.04mol) and the resultant suspension was stirred at 0-5° for a further 2 hours.

The reaction mixture was washed with 2 M-  
20 aqueous hydrochloric acid (1 x 240 mL, 1 x 160 mL) and the combined aqueous acidic washes were back extracted with dichloromethane (160 mL). The organic phases were combined, clarified, and concentrated in *vacuo* to c.a. 240 mL, 2,2,4-trimethylpentane (400 mL) was added and  
25 the solution concentrated, in *vacuo*, to 240 mL.

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Crystallisation of the product occurred during the distillation. Further 2,2,4-trimethylpentane (400 mL) was added and the mixture concentrated to c.a. 700 mL. The stirred suspension was then cooled to 5° and aged  
5 for 60 minutes. The solid was collected by vacuum filtration, washed with 2,2,4-trimethylpentane (2 x 80 mL) and dried, *in vacuo*, at 33° to give the title compound (93.2 g, 68%) as shown by comparison of the <sup>1</sup>H NMR spectrum with that of Example 8.

10           b) Oxalyl chloride (102 g, 0.80 mol) was added over 20 minutes to a stirred, cold (-10°) mixture of N,N-dimethylformamide (63 mL) and dichloromethane (840 mL) and the suspension formed was stirred at -10° to -6° for 15 minutes. The compound of Example B (140  
15 g, 0.728 mol) was added and the resultant pale yellow solution was stirred at -8° for 20 minutes.  
(1R,2S,5R)-(-)-Menthol (126 g, 0.80 mol) was added followed by pyridine (140 mL, 1.73 mol), added over 50 minutes. The suspension formed was stirred at -9° for  
20 18 hours and then 1 M aqueous hydrochloric acid (280 mL) was added. The separated aqueous acid phase was extracted with dichloromethane (140 mL) and the combined organic phases were washed with 1 M aqueous hydrochloric acid (280 mL). The aqueous phase was back  
25 extracted with dichloromethane (140 mL) and the combined organic phases were washed with a solution containing sodium hydrogen carbonate (5.6 g) and sodium chloride (28 g) in water (266 mL). The aqueous phase was back extracted with dichloromethane (140 mL) and  
30 the combined organic phases were clarified and concentrated to 560 mL by distillation at atmospheric pressure. 2,2,4-Trimethylpentane (700 mL) was added and the solution was concentrated, *in vacuo*, to 700 mL. The 2,2,4-trimethylpentane addition/reconcentration  
35 procedure was repeated, and the resultant solution was

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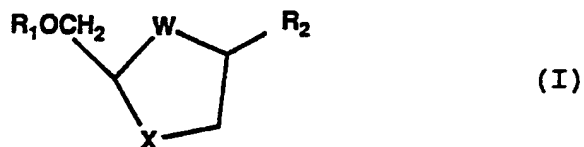
cooled to 17° (seeded with authentic product (0.7 g) at 34° and 23°). The suspension was stirred at 17° for 2 hours and the solid was collected by vacuum filtration, washed with 2,2,4-trimethylpentane (2 x 70 mL) and  
5 dried, *in vacuo*, at 43° to give the title compound (332 g, 14%) as shown by comparison of the <sup>1</sup>H-NMR spectrum with that of Example 8).

While we have presented a number of embodiments of our invention, many alternatives,  
10 modifications and variations of these embodiments will be apparent to those of ordinary skill in the art. Therefore, it will be appreciated that the scope of this invention is to be defined by the following claims, rather than the specific examples presented  
15 above.

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## Claims:

1. A diastereoselective process for producing optically active cis-nucleoside and nucleoside analogues and derivatives of formula (I)



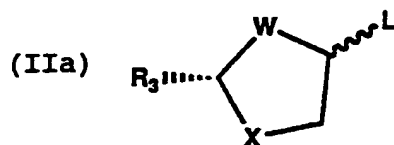
wherein W is S, S=O, SO<sub>2</sub>, or O;

X is S, S=O, SO<sub>2</sub>, or O;

R<sub>1</sub> is hydrogen or acyl; and

R<sub>2</sub> is a desired purine or pyrimidine base or an analogue or derivative thereof

the process comprising the step of glycosylating the desired purine or pyrimidine base or analogue or derivative thereof with an intermediate of formula (IIa) or (IIb)

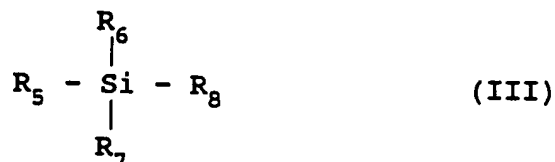


wherein R<sub>3</sub> is a substituted carbonyl or carbonyl derivative; and

L is a leaving group,

using a Lewis acid of the formula (III)

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wherein  $R_5$ ,  $R_6$  and  $R_7$  are independently selected from the group consisting of hydrogen;  $C_{1-20}$  alkyl optionally substituted by fluoro, bromo, chloro, iodo,  $C_{1-6}$  alkoxy or  $C_{6-20}$  aryloxy;  $C_{7-20}$  aralkyl optionally substituted by halogen,  $C_{1-20}$  alkyl or  $C_{1-20}$  alkoxy;  $C_{6-20}$  aryl optionally substituted by fluoro, bromo, chloro, iodo,  $C_{1-20}$  alkyl or  $C_{1-20}$  alkoxy; trialkylsilyl; fluoro; bromo; chloro and iodo; and

$R_8$  is selected from the group consisting of fluoro; bromo; chloro; iodo;  $C_{1-20}$  sulphonate esters, optionally substituted by fluoro, bromo, chloro or iodo;  $C_{1-20}$  alkyl esters optionally substituted by fluoro, bromo, chloro or iodo; polyvalent halides; trisubstituted silyl groups of the general formula  $(R_5)(R_6)(R_7)Si$  (wherein  $R_5$ ,  $R_6$ , and  $R_7$  are as defined above); saturated or unsaturated selenenyl  $C_{6-20}$  aryl; substituted or unsubstituted  $C_{6-20}$  arylsulfenyl; substituted or unsubstituted  $C_{6-20}$  alkoxyalkyl; and trialkylsiloxo.

2. The process according to claim 1, further comprising the step of reducing  $R_3$  of the glycosylated purine or pyrimidine base or analogue or derivative thereof to produce the optically active *cis*-nucleoside or nucleoside analogue or derivative of formula (I).

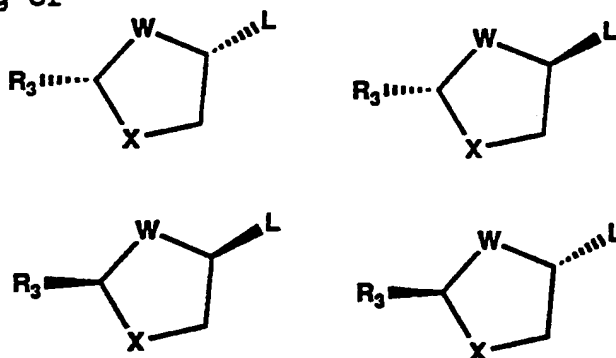
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3. The process according to claim 1, further comprising the step of producing the intermediate of (IIa) or (IIb) by chemically resolving said intermediate from a mixture of (IIa) and (IIb) using a chiral auxiliary.

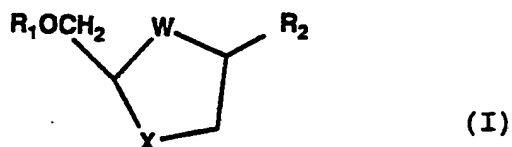
4. The process according to claim 1, wherein the intermediate is the compound of formula (IIa).

5. The process according to claim 1, wherein the intermediate is the compound of formula (IIb).

6. The process according to claim 1, wherein the intermediate is selected from the group consisting of



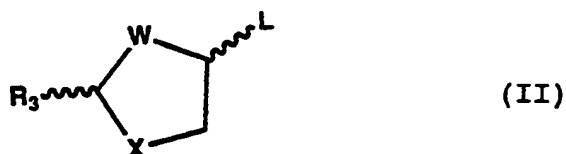
7. A diastereoselective process for producing optically active cis-nucleoside and nucleoside analogues and derivatives of formula (I)



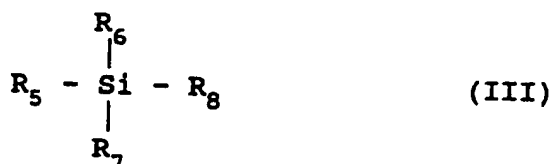
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wherein W is S, S=O, SO<sub>2</sub>, or O;  
 X is S, S=O, SO<sub>2</sub>, or O;  
 R<sub>1</sub> is hydrogen or acyl; and  
 R<sub>2</sub> is a desired purine or pyrimidine base or  
 an analogue or derivative thereof

the process comprising the step of glycosylating the  
 desired purine or pyrimidine base or analogue or  
 derivative thereof with a single enantiomer of a  
 compound of formula (II)



wherein R<sub>3</sub> is a substituted carbonyl or carbonyl  
 derivative; and  
 L is a leaving group,  
 using a Lewis acid of the formula (III)



wherein R<sub>5</sub>, R<sub>6</sub> and R<sub>7</sub> are independently selected from  
 the group consisting of hydrogen; C<sub>1-20</sub> alkyl  
 optionally substituted by fluoro, bromo,  
 chloro, iodo, C<sub>1-6</sub> alkoxy or C<sub>6-20</sub> aryloxy;  
 C<sub>7-20</sub> aralkyl optionally substituted by  
 halogen, C<sub>1-20</sub> alkyl or C<sub>1-20</sub> alkoxy; C<sub>6-20</sub>  
 aryl optionally substituted by fluoro, bromo,  
 chloro, iodo, C<sub>1-20</sub> alkyl or C<sub>1-20</sub> alkoxy;

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trialkylsilyl; fluoro; bromo; chloro and iodo; and

$R_8$  is selected from the group consisting of fluoro; bromo; chloro; iodo;  $C_{1-20}$  sulphonate esters, optionally substituted by fluoro, bromo, chloro or iodo;  $C_{1-20}$  alkyl esters optionally substituted by fluoro, bromo, chloro or iodo; polyvalent halides; trisubstituted silyl groups of the general formula  $(R_5)(R_6)(R_7)Si$  (wherein  $R_5$ ,  $R_6$ , and  $R_7$  are as defined above); saturated or unsaturated selenenyl  $C_{6-20}$  aryl; substituted or unsubstituted  $C_{6-20}$  arylsulfenyl; substituted or unsubstituted  $C_{6-20}$  alkoxyalkyl; and trialkylsiloxo.

8. The process according to claim 7, further comprising the step of reducing  $R_3$  of the glycosylated purine or pyrimidine base or analogue or derivative thereof to produce the optically active *cis*-nucleoside or nucleoside analogue or derivative of formula (I).

9. The process according to claim 7, further comprising the step of resolving the compound of formula (II) into a single enantiomer using a chiral auxiliary before glycosylating the desired purine or pyrimidine base.

10. The process according to any one of claims 1 to 9, wherein W is O and X is S.

11. The process according to claim 10, wherein  $R_2$  is a pyrimidine base.



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12. The process according to claim 11, wherein the pyrimidine base is cytosine or 5-fluorocytosine.

13. The process according to any one of claims 1 to 9, wherein the Lewis acid is selected from the group consisting of trimethylsilyl triflate and iodotrimethylsilane.

14. The process according to claim 3 or 9, wherein the chiral auxiliary is selected from the group consisting of chiral alcohols and chiral amines.

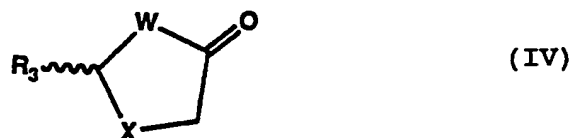
15. The process according to claim 14, wherein the chiral auxiliary is selected from the group consisting of (d)-menthol, (l)-menthol, (+)-norephedrine and (-)-norephedrine.

16. The process according to any one of claims 1 to 9, wherein  $R_3$  is selected from the group consisting of alkoxycarbonyls, carboxyls, diethylcarboxamide, pyrrolidine amide, methyl ketone and phenyl ketone.

17. The process according to claim 16, wherein the  $R_3$  is selected from the group consisting of alkoxycarbonyls and carboxyls.

18. The process according to claim 7 or 8, further comprising the step of producing the compound of formula (II) by chemoselectively reducing the compound of formula (IV)

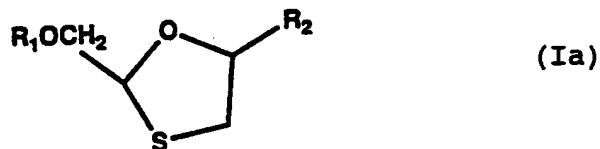
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and converting the resulting hydroxyl group to leaving group L.

19. The process according to claim 18, further comprising the step of reacting the compound of formula (IV) with a chiral auxiliary before it is chemoselectively reduced.

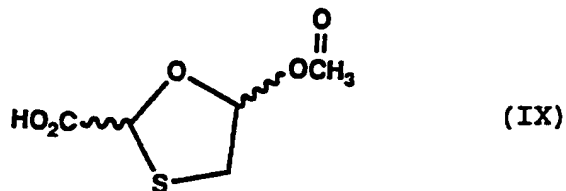
20. A diastereoselective process for producing optically active cis-oxathiolane and analogues and derivatives of formula (Ia)



wherein  $R_1$  is hydrogen or acyl; and

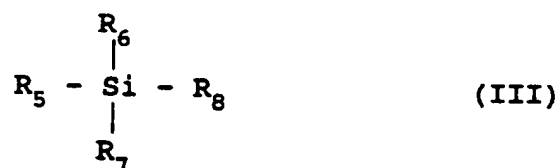
$R_2$  is a desired purine or pyrimidine base or an analogue or derivative thereof

comprising the step of glycosylating the desired purine or pyrimidine base or analogue or derivative thereof with a single enantiomer of a resolved ester derived from the compound of formula (IX)



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using a Lewis acid of the formula (III)



wherein  $R_5$ ,  $R_6$  and  $R_7$  are independently selected from the group consisting of hydrogen;  $C_{1-20}$  alkyl optionally substituted by fluoro, bromo, chloro, iodo,  $C_{1-6}$  alkoxy or  $C_{6-20}$  aryloxy;  $C_{7-20}$  aralkyl optionally substituted by halogen,  $C_{1-20}$  alkyl or  $C_{1-20}$  alkoxy;  $C_{6-20}$  aryl optionally substituted by fluoro, bromo, chloro, iodo,  $C_{1-20}$  alkyl or  $C_{1-20}$  alkoxy; trialkylsilyl; fluoro; bromo; chloro and iodo; and

$R_8$  is selected from the group consisting of fluoro; bromo; chloro; iodo;  $C_{1-20}$  sulphonate esters, optionally substituted by fluoro, bromo, chloro or iodo;  $C_{1-20}$  alkyl esters optionally substituted by fluoro, bromo, chloro or iodo; polyvalent halides; trisubstituted silyl groups of the general formula  $(R_5)(R_6)(R_7)Si$  (wherein  $R_5$ ,  $R_6$ , and  $R_7$  are as defined above); saturated or unsaturated selenenyl  $C_{6-20}$  aryl; substituted or unsubstituted  $C_{6-20}$  arylsulfenyl; substituted or unsubstituted  $C_{6-20}$  alkoxyalkyl; and trialkylsiloxo.

21. The process according to claim 20, further comprising the step of reducing the glycosylated purine or pyrimidine base or analogue or

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derivative thereof to produce the optically active *cis*-oxathiolane or analogue or derivative of formula (Ia).

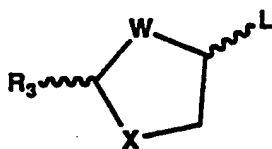
22. The process according to claim 20, further comprising the step of producing the single enantiomer of the ester derived from formula (IX) using a chiral auxiliary.

23. The process according to claim 22, wherein the chiral auxiliary is selected from the group consisting of (d)-menthol and (l)-menthol.

24. The process according to claim 20, wherein  $R_2$  is a pyrimidine base.

25. The process according to claim 24, wherein the pyrimidine base is cytosine or 5-fluorocytosine.

26. An intermediate of formula (II)



(II)

wherein W is S, S=O, SO<sub>2</sub>, or O;

X is S, S=O, SO<sub>2</sub>, or O;

$R_3$  is a substituted carbonyl or carbonyl derivative; and

L is a leaving group.

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27. An intermediate of formula (IIa)



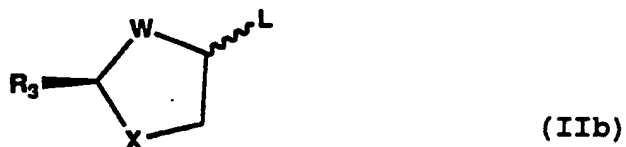
wherein W is S, S=O, SO<sub>2</sub>, or O;

X is S, S=O, SO<sub>2</sub>, or O;

R<sub>3</sub> is a substituted carbonyl or carbonyl derivative; and

L is a leaving group.

28. An intermediate of formula (IIb)



wherein W is S, S=O, SO<sub>2</sub>, or O;

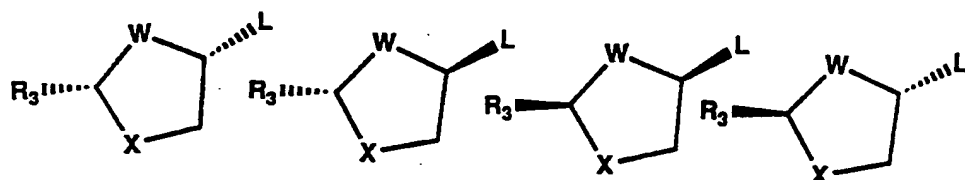
X is S, S=O, SO<sub>2</sub>, or O;

R<sub>3</sub> is a substituted carbonyl or carbonyl derivative; and

L is a leaving group.

29. An intermediate according to claim 26 selected from the group consisting of

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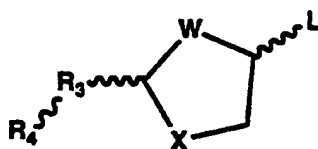
wherein W is S, S=O, SO<sub>2</sub>, or O;

X is S, S=O, SO<sub>2</sub>, or O;

R<sub>3</sub> is a substituted carbonyl or carbonyl derivative; and

L is a leaving group.

30. An intermediate of formula (VI)



(VI)

wherein W is S, S=O, SO<sub>2</sub>, or O;

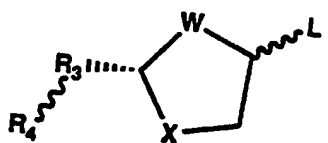
X is S, S=O, SO<sub>2</sub>, or O;

R<sub>3</sub> is a substituted carbonyl or carbonyl derivative;

R<sub>4</sub> is a chiral auxiliary; and

L is a leaving group.

31. An intermediate of formula (VIa)



(VIa)

wherein W is S, S=O, SO<sub>2</sub>, or O;

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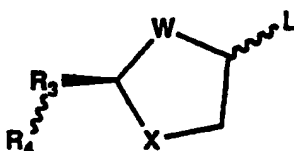
X is S, S=O, SO<sub>2</sub>, or O;

R<sub>3</sub> is a substituted carbonyl or carbonyl derivative;

R<sub>4</sub> is a chiral auxiliary; and

L is a leaving group.

32. An intermediate of formula (VIb)



(VIb)

wherein W is S, S=O, SO<sub>2</sub>, or O;

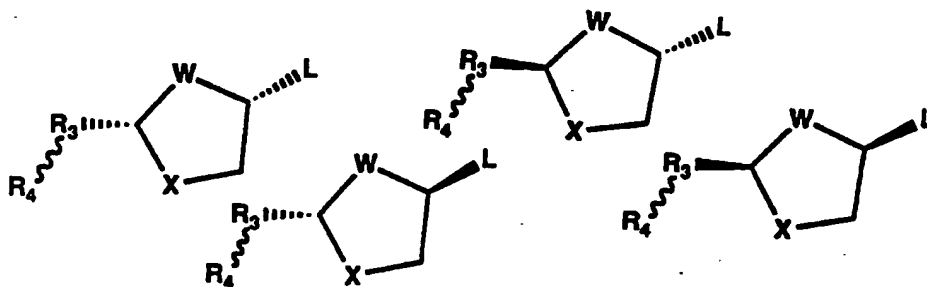
X is S, S=O, SO<sub>2</sub>, or O;

R<sub>3</sub> is a substituted carbonyl or carbonyl derivative;

R<sub>4</sub> is a chiral auxiliary; and

L is a leaving group.

33. An intermediate according to claim 30 selected from the group consisting of



wherein W is S, S=O, SO<sub>2</sub>, or O;

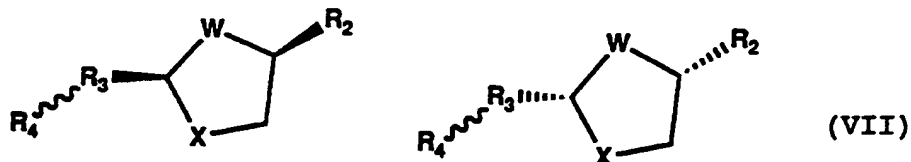
X is S, S=O, SO<sub>2</sub>, or O;

R<sub>3</sub> is a substituted carbonyl or carbonyl derivative;

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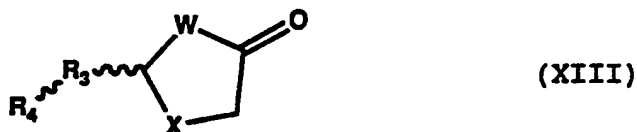
$R_4$  is a chiral auxiliary; and  
 $L$  is a leaving group.

34. An intermediate of formula (VII)



wherein  $W$  is S, S=O, SO<sub>2</sub>, or O;  
 $X$  is S, S=O, SO<sub>2</sub>, or O;  
 $R_2$  is a purine or pyrimidine base or an  
analogue or derivative thereof;  
 $R_3$  is a substituted carbonyl or carbonyl  
derivative; and  
 $R_4$  is a chiral auxiliary.

35. An intermediate of formula (XIII)



wherein  $W$  is S, S=O, SO<sub>2</sub>, or O;  
 $X$  is S, S=O, SO<sub>2</sub>, or O;  
 $R_3$  is a substituted carbonyl or carbonyl  
derivative; and  
 $R_4$  is a chiral auxiliary.



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36. An intermediate according to any one of claims 30 to 35 wherein  $R_4$  is selected from the group consisting of (d)-menthol and (l)-menthol.

37. An intermediate of formula (XIV)



wherein W is S, S=O, SO<sub>2</sub>, or O;

X is S, S=O, SO<sub>2</sub>, or O;

$R_3$  is a substituted carbonyl or carbonyl derivative.

38. An intermediate selected from the group consisting of

- trans*-5-hydroxyoxathiolane-2-carboxylic acid;
- (1'R,2'S,5'R)-menthyl-1,3-oxathiolan-5-one-2S-carboxylate;
- (1'R,2'S,5'R)-menthyl-1,3-oxathiolan-5-one-2R-carboxylate;
- (1'R,2'S,5'R)-menthyl-5S-hydroxy-1,3-oxathiolane-2S-carboxylate;
- (1'R,2'S,5'R)-menthyl-5R-hydroxy-1,3-oxathiolane-2R-carboxylate;
- (1'R,2'S,5'R)-menthyl-5S-hydroxy-1,3-oxathiolane-2R-carboxylate;
- (1'R,2'S,5'R)-menthyl-5R-hydroxy-1,3-oxathiolane-2S-carboxylate;
- (1'R,2'S,5'R)-menthyl-5S-acetoxy-1,3-oxathiolane-2S-carboxylate;

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(1'R,2'S,5'R)-menthyl-5R-acetoxy-1,3-oxathiolane-2R-carboxylate;

(1'R,2'S,5'R)-menthyl-5S-acetoxy-1,3-oxathiolane-2R-carboxylate;

(1'R,2'S,5'R)-menthyl-5R-acetoxy-1,3-oxathiolane-2S-carboxylate;

(1'S,2'R,5'S)-menthyl-5R-acetoxy-1,3-oxathiolane-2S-carboxylate;

(1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolane-2R-carboxylate;

(1'S,2'R,5'S)-menthyl-5R-acetoxy-1,3-oxathiolane-2R-carboxylate;

(1'S,2'R,5'S)-menthyl-5S-acetoxy-1,3-oxathiolane-2S-carboxylate;

(1'R,2'S,5'R)-menthyl-5S-(cytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate;

(1'S,2'R,5'S)-menthyl-5S-(cytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate;

(1'R,2'S,5'R)-menthyl-5R-(cytosin-1"-yl)-1,3-oxathiolane-2S-carboxylate;

(1'S,2'R,5'S)-menthyl-5R-(cytosin-1"-yl)-1,3-oxathiolane-2S-carboxylate;

(1'R,2'S,5'R)-menthyl-5R-(5"-fluorocytosin-1"-yl)-1,3-oxathiolane-2S-carboxylate;

(1'S,2'R,5'S)-menthyl-5S-(5"-fluorocytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate;

(1'S,2'R,5'S)-menthyl-5S-(N-4"-acetylcytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate;

(1'R,2'S,5'R)-menthyl-5S-(cytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate;

(1'S,2'R,5'S)-menthyl-5S-(N-4"-acetylcytosin-1"-yl)-1,3-oxathiolane-2R-carboxylate;

(1'S,2'R,5'S)-menthyl-1,3-oxathiolane-2R-carboxylate;

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(1'S,2'R,5'S)-menthyl-4R-hydroxy-1,3-oxathiolane-2R-carboxylate and (1'S,2'R,5'S)-menthyl-4S-hydroxy-1,3-oxathiolane-2R-carboxylate;

(1'S,2'R,5'S)-menthyl-4R-chloro-1,3-oxathiolane-2R-carboxylate and (1'S,2'R,5'S)-menthyl-4S-chloro-1,3-oxathiolane-2R-carboxylate;

*cis*-2(N-methyl-N-methoxyaminocarbonyl)-5-(uracil-1'-yl)-1,3-oxathiolane;

*cis*- and *trans*-2-benzoyl-5-acetoxy-1,3-oxathiolane;

*cis*-2-(1'-pyrrolidinocarbonyl)-5-acetoxy-1,3-oxathiolane;

*cis*-2-carbomethoxy-5-(5'-bromouracil-1'-yl)-1,3-oxathiolane;

*cis*-2-carboxyl-5-(uracil-1'-yl)-1,3-oxathiolane;

*cis*-2-(1'-pyrrolidinocarbonyl)-5-(uracil-1'-yl)-1,3-oxathiolane;

*cis* 2-benzoyl-5-(uracil-1'-yl)-1,3-oxathiolane;

*cis*- and *trans*-isopropyl 5-acetoxy-1,3-oxathiolane-2-carboxylate;

*cis*-isopropyl-5-(cytosin-1'-yl)-1,3-oxathiolane-2-carboxylate;

*cis*- and *trans*-*t*-butyl 5-acetoxy-1,3-oxathiolane-2-carboxylate;

*cis*-*t*-butyl-5-(cytosin-1'-yl)-1,3-oxathiolane-2-carboxylate;

*cis*- and *trans*-2-N,N-diethylamidocarbonyl- 5-acetoxy-1,3-oxathiolane;

*cis*-2-N,N-diethylamidocarbonyl-5-(cytosin-1'-yl)-1,3 -oxathiolane;

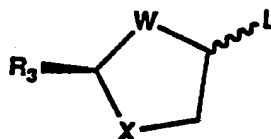
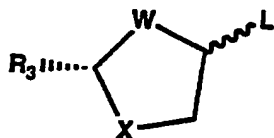
*cis*- and *trans*-2-carboethoxy-4-acetoxy-1,3-dioxolane;

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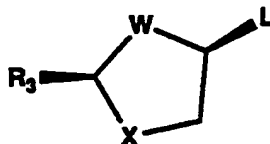
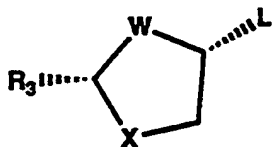
*cis*- and *trans*-2-carboethoxy-4-(thymine-1'-yl)-1,3-dioxolane; and

*cis*- and *trans*-2-carboethoxy-4-(N-4'-acetylcytosine-1'-yl)-1,3-dioxolane.

39. A process for producing an intermediate of formula (IIa) or formula (IIb) comprising the step of resolving a mixture of (IIa) or (IIb) using a chiral auxiliary:



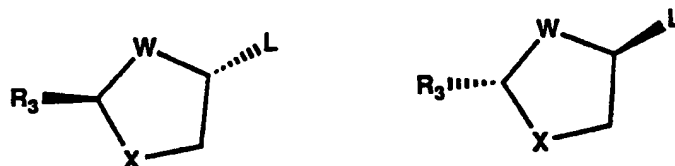
40. A method of preparing a compound of formula



comprising the step of resolving a mixture of the two compounds by means of a chiral auxiliary.

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41. A method of preparing a compound of formula



comprising the step of resolving a mixture of the two compounds by means of a chiral auxiliary.

<b>I. CLASSIFICATION OF SUBJECT MATTER</b> (If several classification symbols apply, indicate all) <sup>6</sup>		
According to International Patent Classification (IPC) or to both National Classification and IPC		
Int.Cl. 5	C07D327/04; C07D411/04;	C07D317/24; C07D411/12; C07D317/34; C07D411/14; C07D405/04 C07D473/00
<b>II. FIELDS SEARCHED</b>		
Minimum Documentation Searched <sup>7</sup>		
Classification System	Classification Symbols	
Int.Cl. 5	C07H ; C07B ; C07D	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched <sup>8</sup>		
<b>III. DOCUMENTS CONSIDERED TO BE RELEVANT<sup>9</sup></b>		
Category <sup>10</sup>	Citation of Document, <sup>11</sup> with indication, where appropriate, of the relevant passages <sup>12</sup>	Relevant to Claim No. <sup>13</sup>
P,A	SYNTHESIS. no. 11, November 1991, STUTTGART DE pages 1046 - 1048; J-L KRAUS ET AL: 'Synthesis of New 2,5-Substituted 1,3-Oxathiolanes. Intermediates in Nucleoside Chemistry'	1
P,X	see the whole document ---	26-29
A	JOURNAL OF ORGANIC CHEMISTRY. vol. 46, 1981, EASTON US pages 3353 - 3354; E.VEDEJS ET AL.: 'Method for Sulfide S-Benzylolation or S-Allylation Using Trimethylsilyl Triflate Activated Benzyl or Allyl Ethers' *page 3354, table 1, compounds 7,8,9* ---	26
A	FR,A,1 445 013 (DR KARL THOMAE GMBH) 31 May 1966 see the whole document --- -/-	26
<p><sup>10</sup> Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"&amp;" document member of the same patent family</p>		
<b>IV. CERTIFICATION</b>		
Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report	
17 AUGUST 1992	17. 09. 92	
International Searching Authority	Signature of Authorized Officer	
EUROPEAN PATENT OFFICE	SCOTT J.R. <i>J.R.M. Scott</i>	

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		Relevant to Claim No.
Category <sup>a</sup>	Citation of Document, with indication, where appropriate, of the relevant passages	
P,A	WO,A,9 117 159 (IAF BIOCHEM INTERNATIONAL INC.) 14 November 1991	1
P,X	see page 1, line 1 - page 3, line 37 ---	26-29
A	EP,A,0 382 526 (IAF BIOCHEM INTERNATIONAL INC.) 16 August 1990	1
X	see the whole document ---	26-29
A	EP,A,0 337 713 (IAF BIOCHEM INTERNATIONAL INC.) 18 October 1989	1
X	see the whole document ---	26-29
A	EP,A,0 266 042 (UNIVERSITY OF STRATHCLYDE) 4 May 1988 see abstract see page 3, line 42 - line 51 ---	1

**ANNEX TO THE INTERNATIONAL SEARCH REPORT  
ON INTERNATIONAL PATENT APPLICATION NO. CA 9200211  
SA 59285**

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office EDP file on  
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information. 17/08/92

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